

This Page Is Inserted by IFW Operations
and is not a part of the Official Record

BEST AVAILABLE IMAGES

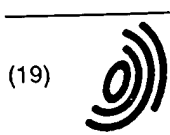
Defective images within this document are accurate representations of the original documents submitted by the applicant.

Defects in the images may include (but are not limited to):

- BLACK BORDERS
- TEXT CUT OFF AT TOP, BOTTOM OR SIDES
- FADED TEXT
- ILLEGIBLE TEXT
- SKEWED/SLANTED IMAGES
- COLORED PHOTOS
- BLACK OR VERY BLACK AND WHITE DARK PHOTOS
- GRAY SCALE DOCUMENTS

IMAGES ARE BEST AVAILABLE COPY.

As rescanning documents *will not* correct images,
please do not report the images to the
Image Problem Mailbox.



Europäisches Patentamt
European Patent Office
Office européen des brevets



(11) EP 1 076 094 A2

(12)

EUROPEAN PATENT APPLICATION

(43) Date of publication:
14.02.2001 Bulletin 2001/07

(51) Int. Cl. 7: C12N 15/74

(21) Application number: 00117225.3

(22) Date of filing: 11.08.2000

(84) Designated Contracting States:
AT BE CH CY DE DK ES FI FR GB GR IE IT LI LU
MC NL PT SE
Designated Extension States:
AL LT LV MK RO SI

(30) Priority: 12.08.1999 JP 22839199

(71) Applicant: Ajinomoto Co., Inc.
Tokyo 104 (JP)

(72) Inventors:
• Matsuzaki, Yumi,
c/o Ajinomoto Co., Inc.
Kawasaki-shi, Kanagawa (JP)
• Kimura, Eiichiro,
c/o Ajinomoto Co., Inc.
Kawasaki-shi, Kanagawa (JP)

- Nakamatsu, Tsuyoshi,
c/o Ajinomoto Co., Inc.
Kawasaki-shi, Kanagawa (JP)
- Kurahashi, Osamu,
c/o Ajinomoto Co., Inc.
Kawasaki-shi, Kanagawa (JP)
- Kawahara, Yoshio,
c/o Ajinomoto Co., Inc.
Kawasaki-shi, Kanagawa (JP)
- Sugimoto, Shinichi,
c/o Ajinomoto Co., Inc.
Kawasaki-shi, Kanagawa (JP)

(74) Representative: HOFFMANN - EITLE
Patent- und Rechtsanwälte
Arabellastrasse 4
81925 München (DE)

(54) Plasmid capable of autonomous replication in coryneform bacteria

(57) A plasmid isolable from *Corynebacterium thermoaminogenes*, which comprises a gene coding for a Rep protein having the amino acid sequence shown in SEQ ID NO: 2 or an amino acid sequence having homology of 90% or more to the amino acid sequence shown in SEQ ID NO: 2, and has a size of about 4.4 kb or about 6 kb, or a derivative thereof.

EP 1 076 094 A2

Description

BACKGROUND OF THE INVENTION

5 [0001] The present invention relates to a novel plasmid derived from *Corynebacterium thermoaminogenes*. The plasmid of the present invention can be utilized for improving of coryneform bacteria, which are used as bacteria for producing useful substances such as L-amino acids.

[0002] Amino acids including L-glutamic acid and L-lysine are produced by fermentative methods using the so-called coryneform bacteria, which generally belong to the genus *Brevibacterium*, *Corynebacterium* or *Microbacterium*, or variant strains thereof (Amino Acid Fermentation, Gakkai Shuppan Center, pp.195-215, 1986).

10 [0003] In the industrial fermentative production of amino acids, besides improvement in yield relative to saccharides, shortening of culture time, improvement in amino acid accumulation concentration and so forth, use of an elevated culture temperature is considered important as a technical factor that raises economical efficiency. That is, culture is usually performed at optimum fermentation temperature, and the optimum temperature is 31.5°C for *Corynebacterium glutamicum*. After the culture is started, heat is generated during the fermentation, and hence amino acid production is markedly reduced if this heat output is not removed. Therefore, cooling equipment is required in order to maintain the temperature of the culture broth to be optimum. On the other hand, if the culture temperature can be elevated, it becomes possible to decrease energy required for cooling and the cooling equipment can be made small.

[0004] Among coryneform bacteria, *Corynebacterium thermoaminogenes* has been isolated as a coryneform bacterium that can grow in a high temperature region (Japanese Patent Application Laid-open (Kokai) No. 63-240779). Whereas growth of *Corynebacterium glutamicum* is markedly suppressed at 40°C, *Corynebacterium thermoaminogenes* can grow at a temperature of about 40°C or higher, and is considered to be suitable for high temperature fermentation.

25 [0005] Currently, improving relying on DNA recombination techniques is progressing in *Escherichia coli* or coryneform bacteria. In order to improve microorganisms by DNA recombination techniques, even plasmids derived from microorganisms belonging to another species or genus or broad host spectrum vectors are often used. However, plasmids proper to objective microorganisms of improving are generally used. In particular, when optimum culture temperature for the objective microorganism of the improving is different from that of microorganisms of the same species or genus, it is preferable to use a plasmid proper to the microorganism.

30 [0006] So far obtained as plasmids derived from coryneform bacteria are pAM330 from *Brevibacterium lactofermentum* ATCC13869 (Japanese Patent Application Laid-open (Kokai) No. 58-67669), pBL1 from *Brevibacterium lactofermentum* ATCC21798 (Santamaria. R. et al., J. Gen. Microbiol., 130, pp.2237-2246, 1984), pHM1519 from *Corynebacterium glutamicum* ATCC13058 (Japanese Patent Application Laid-open (Kokai) No. 58-77895), pCG1 from *Corynebacterium glutamicum* ATCC31808 (Japanese Patent Application Laid-open (Kokai) No. 57-134500) and pGA1 from *Corynebacterium glutamicum* DSM58 (Japanese Patent Application Laid-open (Kokai) No. 9-2603011).

35 [0007] However, no plasmid proper to *Corynebacterium thermoaminogenes* has obtained at present.

SUMMARY OF THE INVENTION

40 [0008] An object of the present invention is to provide a plasmid useful for improving of the coryneform bacterium that can grow at an elevated temperature, *Corynebacterium thermoaminogenes*.

[0009] The inventors of the present invention found that *Corynebacterium thermoaminogenes* AJ12340 (FERM BP-1539), AJ12308 (FERM BP-1540), AJ12309 (FERM BP-1541) and AJ12310 (FERM BP-1542) each harbored a cryptic plasmid proper to each strain, and successfully isolated and identified each plasmid. Thus, they accomplished the present invention.

45 [0010] That is, the present invention provides a plasmid isolable from *Corynebacterium thermoaminogenes*, which comprises a gene (*rep* gene) coding for a Rep protein having the amino acid sequence shown in SEQ ID NO: 2 or an amino acid sequence having homology of 90% or more to the foregoing amino acid sequence, and has a size of about 4.4 kb or about 6 kb, or a derivative thereof.

50 [0011] Examples of the aforementioned plasmid include a plasmid isolable from *Corynebacterium thermoaminogenes* AJ12340 (FERM BP-1539), AJ12308 (FERM BP-1540) or AJ12310 (FERM BP-1542), which has a size of about 4.4 kb and is represented by the restriction map shown in Fig. 1, and a plasmid isolable from *Corynebacterium thermoaminogenes* AJ12309 (FERM BP-1541), which has a size of about 6 kb and is represented by the restriction map shown in Fig. 2.

55 [0012] Specific examples of the aforementioned plasmid include a plasmid which comprises a gene coding for a Rep protein having the amino acid sequence shown in SEQ ID NO: 2, 4 or 6, and a plasmid which comprises a gene coding for a Rep protein having the amino acid sequence shown in SEQ ID NO: 8.

BRIEF EXPLANATION OF THE DRAWINGS

[0013]

- 5 Fig. 1 is a restriction map of the plasmids pYM1, pYM2 and pYM3 of the present invention.
 Fig. 2 is a restriction map of the plasmid pYM4 of the present invention.
 Fig. 3 shows construction of pYMFK.
 Fig. 4 shows construction of pYMK.
 Fig. 5 shows construction of pYMC.
 10 Fig. 6 shows construction of pK1.

DETAILED DESCRIPTION OF THE INVENTION

[0014] The plasmid of the present invention can be isolated from *Corynebacterium thermoaminogenes* AJ12340 (FERM BP-1539), AJ12308 (FERM BP-1540), AJ12309 (FERM BP-1541) or AJ12310 (FERM BP-1542) according to a usual method for preparing a plasmid such as the alkali method (Text for Bioengineering Experiments, Edited by the Society for Bioscience and Bioengineering, Japan, p.105, Baifukan, 1992). As for FERM BP-1539, its original deposition, which was deposited at the National Institute of Bioscience and Human-Technology, Agency of Industrial Science and Technology (postal code 305-8566, 1-3 Higashi 1-chome, Tsukuba-shi, Ibaraki-ken, Japan) on March 13, 1987 and given an accession number of FERM P-9277, was transferred to an international deposition under the provisions of the Budapest Treaty on October 27, 1987 and has been deposited at the same depository. As for FERM BP-1540, FERM BP-1541 and FERM BP-1542, their original depositions, which were deposited at the aforementioned depository on March 10, 1987 and given accession numbers of FERM P-9244, FERM P-9245 and FERM P-9246, were transferred to international depositions under the provisions of the Budapest Treaty on October 27, 1987 and have been deposited at the same depository.

[0015] The inventors of the present invention isolated and identified plasmids each proper to each of the aforementioned *Corynebacterium thermoaminogenes* AJ12308 (FERM BP-1540), AJ12310 (FERM BP-1542), AJ12340 (FERM BP-1539) and AJ12309 (FERM BP-1541) from them, and designated as pYM1, pYM2, pYM3 and pYM4 in that order. These plasmids are plasmids that exist as double-stranded circular DNA in a cell of *Corynebacterium thermoaminogenes*. The nucleotide sequence of the *rep* gene contained in pYM1 is shown in SEQ ID NO: 1, the nucleotide sequence of the *rep* gene contained in pYM2 is shown in SEQ ID NO: 3, the nucleotide sequence of the *rep* gene contained in pYM3 is shown in SEQ ID NO: 5, and the nucleotide sequence of the *rep* gene contained in pYM4 is shown in SEQ ID NO: 7. The amino acid sequences that can be encoded by the *rep* genes contained in these plasmids are shown in SEQ ID NOS: 2, 4, 6 and 8. pYM1, pYM2 and pYM3 each have a size of about 4.4 kb. pYM4 has a size of about 6 kb.

[0016] Numbers and sizes of fragments that can be obtained when pYM1, pYM2 and pYM3 are digested with typical restriction enzymes are shown in Table 1. Numbers and sizes of fragments that can be obtained when pYM4 is digested with typical restriction enzymes are shown in Table 2. Further, a restriction map of pYM1, pYM2 and pYM3 is shown in Fig. 1, and a restriction map of pYM4 is shown in Fig. 2.

Table 1

Restriction enzyme	Number of digestion site	DNA fragment (kb)
<i>Bgl</i> II	0	-
<i>Bam</i> HI	2	1.8, 2.6
<i>Bst</i> PI	1	4.4
<i>Eco</i> RI	1	4.4
<i>Hinc</i> II	4	0.3, 0.5, 2.0, 1.6
<i>Hind</i> III	0	-
<i>Kpn</i> I	0	-
<i>Nae</i> I	2	0.1, 4.3
<i>Nco</i> I	1	4.4
<i>Nhe</i> I	2	1.8, 2.6

Table 1 (continued)

Restriction enzyme	Number of digestion site	DNA fragment (kb)
<i>Pma</i> CI	1	4.4
<i>Sac</i> I	0	-
<i>Sal</i> I	0	-
<i>Sac</i> II	3	0.1, 1.4, 2.9
<i>Sma</i> I	3	0.1, 1.8, 2.5
<i>Sph</i> I	0	-
<i>Tth</i> 111I	1	4.4
<i>Xba</i> I	0	-

Table 2

Restriction enzyme	Number of digestion site	DNA fragment (kb)
<i>Bgl</i> II	1	6.0
<i>Bam</i> HI	2	3.8, 2.2
<i>Bst</i> PI	2	1.2, 4.8
<i>Eco</i> RI	1	6.0
<i>Hinc</i> II	4	0.3, 0.4, 1.2, 1.7, 2.4
<i>Hind</i> III	0	-
<i>Kpn</i> I	0	-
<i>Nae</i> I	2	0.1, 5.9
<i>Nco</i> I	3	0.2, 2.8, 3.0
<i>Nhe</i> I	3	0.1, 2.3, 3.6
<i>Pma</i> CI	0	-
<i>Sac</i> I	0	-
<i>Sal</i> I	0	-
<i>Sac</i> II	5	0.1, 0.2, 0.9, 1.8, 3.0
<i>Sma</i> I	2	0.1, 5.9
<i>Sph</i> I	0	-
<i>Tth</i> 111I	0	-
<i>Xba</i> I	0	-

[0017] Determination of the nucleotide sequence of the plasmid of the present invention revealed that pYM1, pYM2, and pYM3 contained 4368 bp, 4369 bp and 4369 bp, respectively, and they had substantially the same structure and showed homology of 99.9% to one another on the nucleotide sequence level. Further, pYM4 contained 5967 bp and it showed extremely high homology to pYM1, pYM2 and pYM3 for the region of about 4.4 kb except for the region of about 1.6 kb, while it showed homology of about 81% to them as a whole.

[0018] The plasmids contain respective *rep* genes which show high homology to one another. Homology was compared for the amino acid sequences of the Rep proteins encoded by the *rep* genes (SEQ ID NOS: 2, 4, 6 and 8) and the amino acid sequences of the Rep proteins encoded by *rep* genes of known plasmids derived from coryneform bacteria. Homology of 99% or more was observed among pYM1, pYM2 and pYM3, and homology of 81.91% was observed

between pYM2 and pYM4. On the other hand, they showed no homology to the known plasmid pAM330 of a coryneform bacterium, and they showed homology of 80% or less to pGA1 and pCG1. The results are shown in Table 3. Thus, the plasmid of the present invention and the known plasmids of coryneform bacteria are distinguishable based on the homology of the Rep protein.

5 [0019] The homology is calculated according to the method described in Takashi, K. and Gotoh, O., J. Biochem., 92, 1173-1177 (1984).

Table 3

Homology of amino acid sequences of Rep protein encoded by various plasmids				
	PYM2	pYM4	pGA1	pCG1
PYM2	-	81.91%	68.01%	70.73%
PYM4	-	-	69.39%	70.23%
PGA1	-	-	-	75.31%
PCG1	-	-	-	-

20 [0020] Since the plasmid of the present invention can sufficiently replicate in cells of coryneform bacteria including *Corynebacterium thermoaminogenes*, genetic information of a foreign gene can be expressed in a host microorganism by inserting the foreign gene at any site of the plasmid or the derivative thereof, and transforming the host microorganism with the obtained recombinant plasmid.

25 [0021] Examples of coryneform bacteria are listed below.

Corynebacterium acetoacidophilum
Corynebacterium acetoglutamicum
Corynebacterium callunae
30 *Corynebacterium glutamicum*
Corynebacterium thermoaminogenes
Corynebacterium lilium (*Corynebacterium glutamicum*)
Corynebacterium melassecola
Brevibacterium divaricatum (*Corynebacterium glutamicum*)
35 *Brevibacterium lactofermentum* (*Corynebacterium glutamicum*)
Brevibacterium saccharolyticum
Brevibacterium immariophilum
Brevibacterium roseum
Brevibacterium flavum (*Corynebacterium glutamicum*)
40 *Brevibacterium thiogenitalis*

[0022] A derivative of the plasmid of the present invention means a plasmid composed of a part of the plasmid of the present invention, or a part of the plasmid of the present invention or the plasmid of present invention and another DNA sequence. The part means a part containing a region essential for the autonomous replication of the plasmid. The plasmid of the present invention can replicate in a host microorganism even if a region other than the region essential for the autonomous replication of the plasmid (replication control region), that is, the region other than the region containing the replication origin and genes necessary for the replication, is deleted. In addition, a plasmid including such a deletion has a smaller size. Therefore, a plasmid having such a deletion is preferred for use as a vector. Further, if a marker gene such as a drug resistance gene is inserted into the plasmid of the present invention or a part thereof, it becomes easy to detect transformants thanks to phenotype of the marker gene in the transformants. Examples of such a marker gene that can be used in the host include a chloramphenicol resistance gene, kanamycin resistance gene, streptomycin resistance gene, tetracycline resistance gene, trimethoprim resistance gene, erythromycin resistance gene and so forth.

55 [0023] Further, if the plasmid of the present invention is made as a shuttle vector autonomously replicable in coryneform bacteria and other bacteria such as *Escherichia coli* by ligating the plasmid of the present invention or a part thereof with a plasmid autonomously replicable in the other bacteria such as *Escherichia coli* or a part thereof containing a replication control region thereof, manipulations such as preparation of plasmid and preparation of recombinant plasmid containing a target gene can be performed using *Escherichia coli*. Examples of the plasmid

autonomously replicable in *Escherichia coli* include, for example, pUC19, pUC18, pBR322, pHSG299, pHSG298, pHSG399, pHSG398, RSF1010, pMW119, pMW118, pMW219, pMW218 and so forth.

[0024] Although pYM1, pYM2, pYM3 and pYM4 themselves are characterized by the restriction maps shown in Figs. 1 and 2, the plasmid of present invention is not necessarily required to have these restriction maps, and any restriction site may be deleted so long as such deletion does not affect the autonomous replication ability. Further, the plasmid of the present invention may contain a restriction site that is not contained in pYM1, pYM2, pYM3 and pYM4.

[0025] The derivative of the plasmid as described above can be constructed in the same manner as the conventionally known construction of cloning vectors, expression vectors and so forth. In order to construct the derivative, it is preferable to determine the nucleotide sequences of pYM1, pYM2, pYM3 and pYM4. The nucleotide sequence can be determined by known methods such as the dideoxy method.

[0026] In order to insert a foreign gene into the plasmid or the derivative thereof of the present invention, it is convenient to insert it into a restriction site of the plasmid or the derivative thereof. As such a restriction site, one present as a single digestion site is preferred. In order to insert a foreign gene, the plasmid and a source of the foreign gene such as genome DNA can be partially or fully digested with one or more restriction enzymes that provide the same cohesive ends for the both, e.g., the same restriction enzyme, and they can be ligated under a suitable condition. They may also be ligated at blunt ends.

[0027] For preparation of plasmid DNA, digestion and ligation of DNA, transformation and so forth, those methods well known to those skilled in the art may be employed. Such methods are described in Sambrook, J., Fritsch, E.F., and Maniatis, T., "Molecular Cloning: A Laboratory Manual, Second Edition", Cold Spring Harbor Laboratory Press (1989) and so forth.

[0028] According to the present invention, a novel plasmid derived from *Corynebacterium thermoaminogenes* is provided as described above.

EXAMPLES

[0029] Hereafter, the present invention will be explained in more detail with reference to the following examples.

Example 1

Isolation and characterization of plasmids from *Corynebacterium thermoaminogenes* (FERM BP-1539, FERM BP-1540, FERM BP-1541, FERM BP-1542)

[0030] *Corynebacterium thermoaminogenes* AJ12340 (FERM BP-1539), AJ12308 (FERM BP-1540), AJ12309 (FERM BP-1541) and AJ12310 (FERM BP-1542) were cultured for 12 hours in CM2B liquid medium (Bacto-trypton (Difco): 1%, Bacto-yeast-extract (Difco): 1%, NaCl: 0.5%, biotin: 10 µg/L), and plasmid DNA fractions were obtained by the alkali method (Text for Bioengineering Experiments, Edited by the Society for Bioscience and Bioengineering, Japan, p.105, Baifukan, 1992). When these fractions were analyzed by agarose gel electrophoresis (Sambrook, J., Fritsch, E.F., and Maniatis, T., "Molecular Cloning: A Laboratory Manual, Second Edition", Cold Spring Harbor Laboratory Press (1989)), DNA bands were detected for all of the cases, and hence it was demonstrated that the aforementioned strains harbored plasmids. The plasmids prepared from FERM BP-1540, FERM BP-1542 and FERM BP-1539 were designated as pYM1, pYM2 and pYM3, respectively. The plasmid prepared from FERM BP-1541 was designated as pYM4. The plasmids pYM1, pYM2 and pYM3 each had a length of about 4.4 kb, and the plasmid pYM4 had a length of about 6.0 kb.

[0031] The plasmids pYM1, pYM2, pYM3 and pYM4 were digested with restriction enzymes *Bgl*II, *Bam*HI, *Bst*PI, *Eco*RI, *Hinc*II, *Hind*III, *Kpn*I, *Nae*I, *Nco*I, *Nhe*I, *Pma*CI, *Sac*I, *Sac*II, *Sal*I, *Sma*I, *Sph*I, *Tth*111I and *Xba*I (produced by Takara Co.), and lengths of the produced DNA fragments were measured by agarose gel electrophoresis. The electrophoresis was performed at 100 V/cm and a constant voltage for several hours by using 0.8% agarose gel. As molecular weight markers, λ phage DNA (Takara Shuzo) digested with a restriction enzyme *Hind*III was used. The results obtained for pYM1, pYM2 and pYM3 are shown in Table 1. The results obtained for pYM4 are shown in Table 2. The restriction map of pYM1, pYM2 and pYM3 is shown in Fig. 1, and the restriction map of pYM4 is shown in Fig. 2, which were prepared based on the above results.

[0032] The results of nucleotide sequencing of pYM1, pYM2, pYM3 and pYM4 by the dideoxy method are shown in SEQ ID NOS: 1, 3, 5 and 7 in that order.

Example 2

Construction of shuttle vector pYMFK containing Km resistance gene derived from *Streptococcus faecalis*

- 5 [0033] As a region necessary for efficient replication of pYM2 in coryneform bacteria, there are present an AT-rich region upstream from *rep* and a region affecting copy number downstream from *rep*, besides the region coding for *rep*.
- [0034] Therefore, in order to obtain a shuttle vector that can replicate in coryneform bacteria and *E. coli* without impairing the replication ability of pYM2, a region enabling autonomous replication in *E. coli* and a selection marker were inserted into sites in the vicinity of the *Bst*PI site of pYM2.
- 10 [0035] First, a vector having a drug resistance gene of *S. faecalis* was constructed. The kanamycin resistance gene of *S. faecalis* was amplified by PCR from a known plasmid containing that gene. The nucleotide sequence of the kanamycin resistance gene of the *S. faecalis* has already been elucidated (Trieu-Cuot, P. and Courvalin, P., *Gene*, 23 (3), pp.331-341 (1983)). Based on this sequence, the primers having the nucleotide sequences shown as SEQ ID NOS: 16 and 17 were synthesized, and PCR was performed by using pDG783 (Anne-Marie Guerout-Fleury et al., *Gene*, 167, pp.335-337 (1995)) as a template to amplify a DNA fragment containing the kanamycin resistance gene and its promoter.
- 15 [0036] The above DNA fragment was purified by using SUPREC02 produced by Takara Shuzo Co., Ltd., completely digested with restriction enzymes *Hind*III and *Hinc*II, and blunt-ended. The blunt-ending was performed by using Blunting Kit produced by Takara Shuzo Co., Ltd. This DNA fragment and an amplification product obtained by PCR utilizing the primers having the nucleotide sequences shown as SEQ ID NOS: 18 and 19 and pHSG399 (see S. Takeshita et al., *Gene*, 61, pp.63-74 (1987)) as a template and purification and blunt-ending of the PCR product were mixed and ligated. The ligation reaction was performed by using DNA Ligation Kit ver.2 produced by Takara Shuzo Co., Ltd. Competent cells of *Escherichia coli* JM109 (produced by Takara Shuzo Co., Ltd.) were transformed with the ligated DNA, and applied to L medium (10 g/L of Bacto trypton, 5 g/L of Bacto yeast extract, 5 g/L of NaCl, and 15 g/L of agar, pH 7.2) containing 10 µg/ml of IPTG (isopropyl-β-D-thiogalactopyranoside), 40 µg/ml of X-Gal (5-bromo-4-chloro-3-indolyl-β-D-galactoside) and 25 µg/ml of kanamycin, and cultured overnight. Then, the formed blue colonies were picked up, and subjected to single colony isolation to obtain transformants.
- 20 [0037] Plasmids were prepared from the transformants by using the alkaline method (Text for Bioengineering Experiments, Edited by the Society for Bioscience and Bioengineering, Japan, p.105, Baifukan, 1992), and restriction maps were prepared. A plasmid having a restriction map equivalent to that shown at a lower position in Fig. 6 was designated as pK1. This plasmid is stably harbored in *Escherichia coli*, and imparts kanamycin resistance to a host. Moreover, since it contains the *lacZ'* gene, it is suitable for use as a cloning vector.
- 25 [0038] Then, a region containing the replication origin was amplified by Pyrobest-Taq (Takara Shuzo Co., Ltd.) using pYM2 extracted from *C. thermoaminogenes* AJ12310 (FERM BP-1542) as a template (The entire nucleotide sequence of pYM2 is shown in SEQ ID NO: 9.) and the following primers prepared based on a sequence in pYM2 near the *Bst*PI site:

S1: 5'-AAC CAG GGG GAG GGC GCG AGG C-3' (SEQ ID NO: 10)

S3: 5'-TCT CGT AGG CTG CAT CCG AGG CGG GG-3' (SEQ ID NO: 11)

- 40 The reaction condition was 94°C for 5 minutes, then a cycle of 98°C for 20 seconds and 68°C for 4 minutes, which was repeated for 30 cycles, and 72°C for 4 minutes. After the reaction, the mixture was stored at 4°C.

- [0039] The obtained amplified fragment was purified by using MicroSpin TM S-400 HR columns produced by Amersham Pharmacia Biotech Co., blunt-ended by using DNA Blunting Kit produced by Takara Shuzo Co., Ltd., and then ligated to pK1 treated with *Hinc*II by using DNA Ligation Kit. ver. 2 produced by Takara Shuzo Co., Ltd. Competent cells of *Escherichia coli* JM109 (produced by Takara Shuzo) were transformed with the ligated DNA to obtain transformant strains.

- [0040] Plasmids were prepared from the transformant strains using the alkali method (Text for Bioengineering Experiments, Edited by the Society for Bioscience and Bioengineering, Japan, p.105, Baifukan, 1992) and restriction maps of the plasmids were prepared. One showing a restriction map equivalent to that shown at a lower position in Fig. 3 was designated as pYMFK. pYMFK had a size of about 7.0 kb, and was able to autonomously replicate in *E. coli* and coryneform bacteria and impart Km resistance to a host.

Example 3

- 55 Construction of pYMK containing Km resistance gene derived from Tn903

[0041] A region containing the replication origin was amplified in the same manner as in Example 2 by using pYM2

extracted from *C. thermoaminogenes* AJ12310 (FERM BP-1542) as a template and the following primers:

S1XbaI: 5'-GCT CTA GAG CAA CCA GGG GGA GGG CGC GAG GC-3' (SEQ ID NO: 12)
S3XbaI: 5'-GCT CTA GAG CTC TCG TAG GCT GCA TCG GAG GCG GGG-3' (SEQ ID NO: 13)

5 [0042] The obtained amplified fragment was purified by using MicroSpin TM S-400 HR columns produced by Amersham Pharmacia Biotech Co., digested with a restriction enzyme *XbaI* produced by Takara Shuzo Co., Ltd., and then ligated to a fragment obtained by fully digesting pHSG299 (Takara Shuzo Co., Ltd.) with *XbaI* by using DNA Ligation Kit. ver. 2 produced by Takara Shuzo Co., Ltd. Competent cells of *Escherichia coli* JM109 (produced by Takara Shuzo)

10 were transformed with the ligated DNA to obtain transformant strains.

[0043] Plasmids were prepared from the transformant strains using the alkali method (Text for Bioengineering Experiments, Edited by the Society for Bioscience and Bioengineering, Japan, p.105, Baifukan, 1992) and restriction maps of the plasmids were prepared. One showing a restriction map equivalent to that shown at a lower position in Fig. 4 was designated as pYMK. pYMK had a size of about 7.0 kb, and was able to autonomously replicate in *E. coli* and

15 coryneform bacteria and impart Km resistance to a host.

Example 4

Construction of shuttle vector pYMC containing Cm resistance gene derived from Tn9

20 [0044] A region containing the replication origin was amplified in the same manner as in Example 2 by using pYM2 extracted from *C. thermoaminogenes* AJ12310 (FERM BP-1542) as a template and the following primers:

25 S1XbaI: 5'-GCT CTA GAG CAA CCA GGG GGA GGG CGC GAG GC-3' (SEQ ID NO: 14)
S3XbaI: 5'-GCT CTA GAG CTC TCG TAG GCT GCA TCG GAG GCG GGG-3' (SEQ ID NO: 15)

[0045] The above DNA was purified by using MicroSpin TM S-400 HR columns produced by Amersham Pharmacia Biotech Co., digested with a restriction enzyme *XbaI* produced by Takara Shuzo Co., Ltd., and then ligated to a fragment obtained by treating pHSG399 (Takara Shuzo Co., Ltd.) with *XbaI* by using DNA Ligation Kit. ver. 2 produced by

30 Takara Shuzo Co., Ltd. Competent cells of *Escherichia coli* JM109 (produced by Takara Shuzo) were transformed with the ligated DNA to obtain transformant strains.

[0046] Plasmids were prepared from the transformant strains using the alkali method (Text for Bioengineering Experiments, Edited by the Society for Bioscience and Bioengineering, Japan, p.105, Baifukan, 1992) and restriction maps of the plasmids were prepared. One showing a restriction map equivalent to that shown at a lower position in Fig. 5 was designated as pYMC. pYMC had a size of about 6.6 kb, and was able to autonomously replicate in *E. coli* and

35 coryneform bacteria and impart Cm resistance to a host.

40

45

50

55

SEQUENCE LISTING

5 <110> Ajinomoto Co., Inc

<120> Plasmid Autonomously Replicable in Coryneform Bacteria

10 <130>

<150> JP 11-228391

<151> 1999-08-12

15 <160> 19

<170> PatentIn Ver. 2.0

<210> 1

20 <211> 1479

<212> DNA

<213> *Corynebacterium thermoaminogenes*

<220>

25 <221> CDS

<222> (1)..(1476)

<400> 1

atg act cta gcg gat tcg cca gga aca tac aca gca gat gcg tgg aat	48
Met Thr Leu Ala Asp Ser Pro Gly Thr Tyr Thr Ala Asp Ala Trp Asn	
1 5 10 15	
tac tcc act gat ctg ttc gac acc cac cct gag ctg gct tta cgc tcc	96
Tyr Ser Thr Asp Leu Phe Asp Thr His Pro Glu Leu Ala Leu Arg Ser	
20 25 30	
cgg ggt tgg aat cac cag gac gcc gcc gag ttc ctg gcc cac ctg gat	144
Arg Gly Trp Asn His Gln Asp Ala Ala Glu Phe Leu Ala His Leu Asp	
35 40 45	
cgc agc atg ttt cac ggg tgc ccc acc cgg gat ttc tcc gcg gcc tgg	192
Arg Ser Met Phe His Gly Cys Pro Thr Arg Asp Phe Ser Ala Ala Trp	
50 55 60	
gtc aaa gac ccg gaa acc gga gaa acc cgc ccc aag ctg cac aga gtt	240
Val Lys Asp Pro Glu Thr Gly Glu Thr Arg Pro Lys Leu His Arg Val	
65 70 75 80	
ggc acc cgc tca ctt tcc cgg tgc cag tac gtt gcc ctg acc cac ccg	288
Gly Thr Arg Ser Leu Ser Arg Cys Gln Tyr Val Ala Leu Thr His Pro	
85 90 95	
cag cgc tcc gcg gtg ctg gtc tta gac atc gac atc ccc agc cac cag	336
Gln Arg Ser Ala Val Leu Val Leu Asp Ile Asp Ile Pro Ser His Gln	
100 105 110	
gcc gcc ggg aac atc gag cac ctt cac ccg cag gtg tac gcc acc ttg	384
Ala Gly Gly Asn Ile Glu His Leu His Pro Gln Val Tyr Ala Thr Leu	
115 120 125	
gag cgt tgg gca cgg gtg gag aaa gcg ccg gcc tgg atc ggg gtg aac	432
Glu Arg Trp Ala Arg Val Glu Lys Ala Pro Ala Trp Ile Gly Val Asn	

55

EP 1 076 094 A2

	130	135	140	
	ccg ttg tgc gga aag tgc cag ctc atc tgg tgc att gac ccg gtg ttc			480
5	Pro Leu Ser Gly Lys Cys Gln Leu Ile Trp Cys Ile Asp Pro Val Phe			
	145	150	155	160
	gcc gcc gag ggc acc acc agc tgc aac acc cgc ctg cta gcg gcc acc			528
	Ala Ala Glu Gly Thr Thr Ser Ser Asn Thr Arg Leu Leu Ala Ala Thr			
	165	170	175	
10	acc gag gaa atg acc cgt gtg ttc ggc gct gac cag gca ttt tcc cac			576
	Thr Glu Glu Met Thr Arg Val Phe Gly Ala Asp Gln Ala Phe Ser His			
	180	185	190	
	cgg ctg agc cgg tgg ccg ctg cat gtt tct gat gat ccg acc gcg tac			624
	Arg Leu Ser Arg Trp Pro Leu His Val Ser Asp Asp Pro Thr Ala Tyr			
	195	200	205	
15	tcc tgg cac tgc cag cac aac cga gtc gat att ctt gat gag ctg atg			672
	Ser Trp His Cys Gln His Asn Arg Val Asp Ile Leu Asp Glu Leu Met			
	210	215	220	
	gag gta gcc cgc acg atg acc gga tca aaa aag ccc aga gag cac gct			720
	Glu Val Ala Arg Thr Met Thr Gly Ser Lys Lys Pro Arg Glu His Ala			
	225	230	235	240
20	cac cag gag ttt tcc agc ggt cgg gca cgg atc gaa gcc gcg cgg aaa			768
	His Gln Glu Phe Ser Ser Gly Arg Ala Arg Ile Glu Ala Ala Arg Lys			
	245	250	255	
	gcc acc gca gag gcc aaa gcg ctt gcc gcc ttg gac gcc acg ctg cct			816
	Ala Thr Ala Glu Ala Lys Ala Leu Ala Ala Leu Asp Ala Thr Leu Pro			
	260	265	270	
25	acg gcg ctg gag gca tca ggc gat ctc att gac ggg gtg cgg gtg ttg			864
	Thr Ala Leu Glu Ala Ser Gly Asp Leu Ile Asp Gly Val Arg Val Leu			
	275	280	285	
	tgg gca gca gag ggg cgt gca gcc cgt gat gag aca gcg ttt cgc cat			912
	Trp Ala Ala Glu Gly Arg Ala Ala Arg Asp Glu Thr Ala Phe Arg His			
	290	295	300	
30	gcg ttg acc gtg ggt tat cag ctt aaa gcc gca ggt gaa cgc ctg aaa			960
	Ala Leu Thr Val Gly Tyr Gln Leu Lys Ala Ala Gly Glu Arg Leu Lys			
	305	310	315	320
	gat gcc aag atc att gat gcg tat gag cgt gcc tac aac gtc gcc cag			1008
	Asp Ala Lys Ile Ile Asp Ala Tyr Glu Arg Ala Tyr Asn Val Ala Gln			
	325	330	335	
35	gcg gtg gga gct gat ggg cgt gaa ccg gat ctg cct gcc atg cgt gat			1056
	Ala Val Gly Ala Asp Gly Arg Glu Pro Asp Leu Pro Ala Met Arg Asp			
	340	345	350	
	cgt cag acg atg gcc cgc cgt gtg cgc gcc tac gtc gcc aaa ggc cag			1104
	Arg Gln Thr Met Ala Arg Arg Val Arg Ala Tyr Val Ala Lys Gly Gln			
	355	360	365	
	ccc acg gtc agc gcc agc agc aca cag acc cag agc agt cgg gcc cgg			1152
	Pro Thr Val Ser Ala Arg Ser Thr Gln Thr Gln Ser Ser Arg Gly Arg			
	370	375	380	
40	aaa gcc ctg gcc acc atg ggc cgc aga ggc ggg caa aaa gcc gct gaa			1200
	Lys Ala Leu Ala Thr Met Gly Arg Arg Gly Gly Gln Lys Ala Ala Glu			
	385	390	395	400
	cgc tgg aaa acc gat cct aac ggc aaa tac gcc caa gaa aac cgc caa			1248
	Arg Trp Lys Thr Asp Pro Asn Gly Lys Tyr Ala Gln Glu Asn Arg Gln			
	405	410	415	
45	cga ctc gaa gct gca aac aag cga cgt caa gtc agc tgg aac aaa tac			1296
	Arg Leu Glu Ala Ala Asn Lys Arg Arg Gln Val Ser Trp Asn Lys Tyr			
	420	425	430	
50	gcg agc acg aat tct ggc tac ggt ttc cga cac gta tgg gcc agc ttg			1344
55				

Ala Ser Thr Asn Ser Gly Tyr Gly Phe Arg His Val Trp Ala Ser Leu
 435 440 445
 gaa aaa tgc cta cgc gac gag caa atc atg gaa gaa aca ggg ctt tca 1392
 5 Glu Lys Cys Leu Arg Asp Glu Gln Ile Met Glu Glu Thr Gly Leu Ser
 450 455 460
 cga gct acc gtg acg cgc cat tgg gtg cac tgc gag agg ctg gcc tgc 1440
 Arg Ala Thr Val Thr Arg His Trp Val His Cys Glu Arg Leu Ala Cys
 465 470 475 480
 tgc caa atc ctt agg ggg gct cac gcc gta gac aga taa 1479
 10 Cys Gln Ile Leu Arg Gly Ala His Ala Val Asp Arg
 485 490

<210> 2
 <211> 492
 15 <212> PRT
 <213> Corynebacterium thermoaminogenes

<400> 2
 Met Thr Leu Ala Asp Ser Pro Gly Thr Tyr Thr Ala Asp Ala Trp Asn
 1 5 10 15
 20 Tyr Ser Thr Asp Leu Phe Asp Thr His Pro Glu Leu Ala Leu Arg Ser
 20 25 30
 Arg Gly Trp Asn His Gln Asp Ala Ala Glu Phe Leu Ala His Leu Asp
 35 40 45
 Arg Ser Met Phe His Gly Cys Pro Thr Arg Asp Phe Ser Ala Ala Trp
 50 55 60
 25 Val Lys Asp Pro Glu Thr Gly Glu Thr Arg Pro Lys Leu His Arg Val
 65 70 75 80
 Gly Thr Arg Ser Leu Ser Arg Cys Gln Tyr Val Ala Leu Thr His Pro
 85 90 95
 Gln Arg Ser Ala Val Leu Val Leu Asp Ile Asp Ile Pro Ser His Gln
 100 105 110
 30 Ala Gly Gly Asn Ile Glu His Leu His Pro Gln Val Tyr Ala Thr Leu
 115 120 125
 Glu Arg Trp Ala Arg Val Glu Lys Ala Pro Ala Trp Ile Gly Val Asn
 130 135 140
 35 Pro Leu Ser Gly Lys Cys Gln Leu Ile Trp Cys Ile Asp Pro Val Phe
 145 150 155 160
 Ala Ala Glu Gly Thr Thr Ser Ser Asn Thr Arg Leu Leu Ala Ala Thr
 165 170 175
 Thr Glu Glu Met Thr Arg Val Phe Gly Ala Asp Gln Ala Phe Ser His
 180 185 190
 40 Arg Leu Ser Arg Trp Pro Leu His Val Ser Asp Asp Pro Thr Ala Tyr
 195 200 205
 Ser Trp His Cys Gln His Asn Arg Val Asp Ile Leu Asp Glu Leu Met
 210 215 220
 Glu Val Ala Arg Thr Met Thr Gly Ser Lys Lys Pro Arg Glu His Ala
 225 230 235 240
 45 His Gln Glu Phe Ser Ser Gly Arg Ala Arg Ile Glu Ala Ala Arg Lys
 245 250 255
 Ala Thr Ala Glu Ala Lys Ala Leu Ala Ala Leu Asp Ala Thr Leu Pro
 260 265 270
 Thr Ala Leu Glu Ala Ser Gly Asp Leu Ile Asp Gly Val Arg Val Leu
 275 280 285
 50 Trp Ala Ala Glu Gly Arg Ala Ala Arg Asp Glu Thr Ala Phe Arg His
 290 295 300

55

Ala Leu Thr Val Gly Tyr Gln Leu Lys Ala Ala Gly Glu Arg Leu Lys
 305 310 315 320
 Asp Ala Lys Ile Ile Asp Ala Tyr Glu Arg Ala Tyr Asn Val Ala Gln
 325 330 335
 Ala Val Gly Ala Asp Gly Arg Glu Pro Asp Leu Pro Ala Met Arg Asp
 340 345 350
 Arg Gln Thr Met Ala Arg Arg Val Arg Ala Tyr Val Ala Lys Gly Gln
 355 360 365
 Pro Thr Val Ser Ala Arg Ser Thr Gln Thr Gln Ser Ser Arg Gly Arg
 370 375 380
 Lys Ala Leu Ala Thr Met Gly Arg Arg Gly Gly Gln Lys Ala Ala Glu
 385 390 395 400
 Arg Trp Lys Thr Asp Pro Asn Gly Lys Tyr Ala Gln Glu Asn Arg Gln
 405 410 415
 Arg Leu Glu Ala Ala Asn Lys Arg Arg Gln Val Ser Trp Asn Lys Tyr
 420 425 430
 Ala Ser Thr Asn Ser Gly Tyr Gly Phe Arg His Val Trp Ala Ser Leu
 435 440 445
 Glu Lys Cys Leu Arg Asp Glu Gln Ile Met Glu Glu Thr Gly Leu Ser
 450 455 460
 Arg Ala Thr Val Thr Arg His Trp Val His Cys Glu Arg Leu Ala Cys
 465 470 475 480
 Cys Gln Ile Leu Arg Gly Ala His Ala Val Asp Arg
 485 490

 <210> 3
 <211> 1479
 <212> DNA
 <213> *Corynebacterium thermoaminogenes*

 <220>
 <221> CDS
 <222> (1)..(1476)

 <400> 3
 atg act cta gcg gat tcg cca gga aca tac aca gca gat gcg tgg aat 48
 Met Thr Leu Ala Asp Ser Pro Gly Thr Tyr Thr Ala Asp Ala Trp Asn
 1 5 10 15
 tac tcc act gat ctg ttc gac acc cac cct gag ctg gct tta cgc tcc 96
 Tyr Ser Thr Asp Leu Phe Asp Thr His Pro Glu Leu Ala Leu Arg Ser
 20 25 30
 cgg ggt tgg aat cac cag gac gcc gca gag ttc ctg gcc cac ctg gat 144
 Arg Gly Trp Asn His Gln Asp Ala Ala Glu Phe Leu Ala His Leu Asp
 35 40 45
 cgc agc atg ttt cac ggg tgc ccc acc cgg gat ttc tcc gcg gcc tgg 192
 Arg Ser Met Phe His Gly Cys Pro Thr Arg Asp Phe Ser Ala Ala Trp
 50 55 60
 gtc aaa gac ccg gaa acc gga gaa acc cgc ccc aag ctg cac aga gtt 240
 Val Lys Asp Pro Glu Thr Gly Glu Thr Arg Pro Lys Leu His Arg Val
 65 70 75 80
 ggc acc cgc tca ctt tcc cgg tgc cag tac gtt gcc ctg acc cac ccg 288
 Gly Thr Arg Ser Leu Ser Arg Cys Gln Tyr Val Ala Leu Thr His Pro
 85 90 95
 cag cgc tcc gcg gtg ctg gtc tta gac atc gac atc ccc agc cac cag 336
 Gln Arg Ser Ala Val Leu Val Leu Asp Ile Asp Ile Pro Ser His Gln
 100 105 110

	gcc ggc ggg aac atc gag cac ctt cac ccg cag gtg tac gcc acc ttg	384
	Ala Gly Gly Asn Ile Glu His Leu His Pro Gln Val Tyr Ala Thr Leu	
	115 120 125	
5	gag cgt tgg gca cgg gtg gag aaa gcg ccg gcc tgg atc ggg gtg aac	432
	Glu Arg Trp Ala Arg Val Glu Lys Ala Pro Ala Trp Ile Gly Val Asn	
	130 135 140	
	ccg ttg tgc gga aag tgc cag ctc atc tgg tgc att gac ccg gtg ttc	480
	Pro Leu Ser Gly Lys Cys Gln Leu Ile Trp Cys Ile Asp Pro Val Phe	
	145 150 155 160	
10	gcc gcc gag gcc acc acc agc tgc aac acc cgc ctg cta gcg gcc acc	528
	Ala Ala Glu Gly Thr Thr Ser Ser Asn Thr Arg Leu Leu Ala Ala Thr	
	165 170 175	
	acc gag gaa atg acc cgt gtg ttc gcc gct gac cag gca ttt tcc cac	576
	Thr Glu Glu Met Thr Arg Val Phe Gly Ala Asp Gln Ala Phe Ser His	
	180 185 190	
15	cgg ctg agc cgg tgg ccg ctg cat gtt ttt gat gat ccg acc gcg tac	624
	Arg Leu Ser Arg Trp Pro Leu His Val Phe Asp Asp Pro Thr Ala Tyr	
	195 200 205	
	tcc tgg cac tgc cag cac aac cga gtc gat att ctt gat gag ctg atg	672
	Ser Trp His Cys Gln His Asn Arg Val Asp Ile Leu Asp Glu Leu Met	
	210 215 220	
	gag gta gcc cgc acg atg acc gga tca aaa aag ccg aga aag cac gct	720
	Glu Val Ala Arg Thr Met Thr Gly Ser Lys Lys Pro Arg Lys His Ala	
	225 230 235 240	
	cac cag gag ttt tcc agc ggt cgg gca cgg atc gaa gcc gcg cgg aaa	768
	His Gln Glu Phe Ser Ser Gly Arg Ala Arg Ile Glu Ala Ala Arg Lys	
	245 250 255	
	gcc acc gca gag gcc aaa gcg ctt gcc gcc ttg gac gcc acg ctg cct	816
	Ala Thr Ala Glu Ala Lys Ala Leu Ala Ala Leu Asp Ala Thr Leu Pro	
	260 265 270	
30	acg gcg ctg gag gca tca gcc gat ctc att gac ggg gtg cgg gtg ttg	864
	Thr Ala Leu Glu Ala Ser Gly Asp Leu Ile Asp Gly Val Arg Val Leu	
	275 280 285	
	tgg gca gca gag ggg cgt gca gcc cgt gat gag aca gcg ttt cgc cat	912
	Trp Ala Ala Glu Gly Arg Ala Ala Arg Asp Glu Thr Ala Phe Arg His	
	290 295 300	
35	gcg ttg acc gtg ggt tat cag ctt aaa gcc gca ggt gaa cgc ctg aaa	960
	Ala Leu Thr Val Gly Tyr Gln Leu Lys Ala Ala Gly Glu Arg Leu Lys	
	305 310 315 320	
	gat gcc aag atc att gat gcg tat gag cgt gcc tac aac gtc gcc cag	1008
	Asp Ala Lys Ile Ile Asp Ala Tyr Glu Arg Ala Tyr Asn Val Ala Gln	
	325 330 335	
40	gcg gtg gga gct gat ggg cgt gaa ccg gat ctg cct gcc atg cgt gat	1056
	Ala Val Gly Ala Asp Gly Arg Glu Pro Asp Leu Pro Ala Met Arg Asp	
	340 345 350	
	cgt cag acg atg gcc cgc cgt gtg cgc gcc tac gtc gcc aaa ggc cag	1104
	Arg Gln Thr Met Ala Arg Arg Val Arg Ala Tyr Val Ala Lys Gly Gln	
	355 360 365	
45	ccc acg gtc agc gcc agg agc aca cag acc cag agc agt cgg ggc cgg	1152
	Pro Thr Val Ser Ala Arg Ser Thr Gln Thr Gln Ser Ser Arg Gly Arg	
	370 375 380	
	aaa gcc ctg gcc acc atg ggc cgc aga gcc ggg caa aaa gcc gct gaa	1200
	Lys Ala Leu Ala Thr Met Gly Arg Arg Gly Gly Gln Lys Ala Ala Glu	
	385 390 395 400	
50	cgc tgg aaa acc gat cct aac ggc aaa tac gcc caa gaa aac cgc caa	1248
	Arg Trp Lys Thr Asp Pro Asn Gly Lys Tyr Ala Gln Glu Asn Arg Gln	

5
 10
 15
 405
 410
 415
 1296
 1344
 1392
 1440
 1479
 485
 490

<210> 4
 <211> 492
 <212> PRT
 <213> *Corynebacterium thermoaminogenes*

25
 30
 35
 40
 45
 50
 55
 400> 4
 Met Thr Leu Ala Asp Ser Pro Gly Thr Tyr Thr Ala Asp Ala Trp Asn
 1 5 10 15
 Tyr Ser Thr Asp Leu Phe Asp Thr His Pro Glu Leu Ala Leu Arg Ser
 20 25 30
 Arg Gly Trp Asn His Gln Asp Ala Ala Glu Phe Leu Ala His Leu Asp
 35 40 45
 Arg Ser Met Phe His Gly Cys Pro Thr Arg Asp Phe Ser Ala Ala Trp
 50 55 60
 Val Lys Asp Pro Glu Thr Gly Glu Thr Arg Pro Lys Leu His Arg Val
 65 70 75 80
 Gly Thr Arg Ser Leu Ser Arg Cys Gln Tyr Val Ala Leu Thr His Pro
 85 90 95
 Gln Arg Ser Ala Val Leu Val Leu Asp Ile Asp Ile Pro Ser His Gln
 100 105 110
 Ala Gly Gly Asn Ile Glu His Leu His Pro Gln Val Tyr Ala Thr Leu
 115 120 125
 Glu Arg Trp Ala Arg Val Glu Lys Ala Pro Ala Trp Ile Gly Val Asn
 130 135 140
 Pro Leu Ser Gly Lys Cys Gln Leu Ile Trp Cys Ile Asp Pro Val Phe
 145 150 155 160
 Ala Ala Glu Gly Thr Thr Ser Ser Asn Thr Arg Leu Leu Ala Ala Thr
 165 170 175
 Thr Glu Glu Met Thr Arg Val Phe Gly Ala Asp Gln Ala Phe Ser His
 180 185 190
 Arg Leu Ser Arg Trp Pro Leu His Val Phe Asp Asp Pro Thr Ala Tyr
 195 200 205
 Ser Trp His Cys Gln His Asn Arg Val Asp Ile Leu Asp Glu Leu Met
 210 215 220
 Glu Val Ala Arg Thr Met Thr Gly Ser Lys Lys Pro Arg Lys His Ala
 225 230 235 240
 His Gln Glu Phe Ser Ser Gly Arg Ala Arg Ile Glu Ala Ala Arg Lys
 245 250 255
 Ala Thr Ala Glu Ala Lys Ala Leu Ala Ala Leu Asp Ala Thr Leu Pro

5 260 265 270
 Thr Ala Leu Glu Ala Ser Gly Asp Leu Ile Asp Gly Val Arg Val Leu
 275 280 285
 Trp Ala Ala Glu Gly Arg Ala Ala Arg Asp Glu Thr Ala Phe Arg His
 290 295 300
 Ala Leu Thr Val Gly Tyr Gln Leu Lys Ala Ala Gly Glu Arg Leu Lys
 305 310 315 320
 Asp Ala Lys Ile Ile Asp Ala Tyr Glu Arg Ala Tyr Asn Val Ala Gln
 325 330 335
 10 Ala Val Gly Ala Asp Gly Arg Glu Pro Asp Leu Pro Ala Met Arg Asp
 340 345 350
 Arg Gln Thr Met Ala Arg Arg Val Arg Ala Tyr Val Ala Lys Gly Gln
 355 360 365
 Pro Thr Val Ser Ala Arg Ser Thr Gln Thr Gln Ser Ser Arg Gly Arg
 370 375 380
 15 Lys Ala Leu Ala Thr Met Gly Arg Arg Gly Gly Gln Lys Ala Ala Glu
 385 390 395 400
 Arg Trp Lys Thr Asp Pro Asn Gly Lys Tyr Ala Gln Glu Asn Arg Gln
 405 410 415
 Arg Leu Glu Ala Ala Asn Lys Arg Arg Gln Val Ser Trp Asn Lys Tyr
 420 425 430
 20 Ala Ser Thr Asn Ser Gly Tyr Gly Phe Arg His Val Trp Ala Ser Leu
 435 440 445
 Glu Lys Cys Leu Arg Asp Glu Gln Ile Met Glu Glu Thr Gly Leu Ser
 450 455 460
 25 Arg Ala Thr Val Thr Arg His Trp Val His Cys Glu Arg Leu Ala Cys
 465 470 475 480
 Cys Gln Ile Leu Arg Gly Ala His Ala Val His Arg
 485 490

 30 <210> 5
 <211> 1479
 <212> DNA
 <213> Corynebacterium thermoaminogenes

 35 <220>
 <221> CDS
 <222> (1)..(1476)

 40 <400> 5
 atg act cta gcg gat tcg cca gga aca tac aca gca gat gcg tgg aat 48
 Met Thr Leu Ala Asp Ser Pro Gly Thr Tyr Thr Ala Asp Ala Trp Asn
 1 5 10 15
 tac tcc act gat ctg ttc gac acc cac cct gag ctg gct tta cgc tcc 96
 Tyr Ser Thr Asp Leu Phe Asp Thr His Pro Glu Leu Ala Leu Arg Ser
 20 25 30
 45 cgg ggt tgg aat cac cag gac gcc gag ttc ctg gcc cac ctg gat 144
 Arg Gly Trp Asn His Gln Asp Ala Ala Glu Phe Leu Ala His Leu Asp
 35 40 45
 cgc agc atg ttt cac ggg tgc ccc acc cgg gat ttc tcc gcg gcc tgg 192
 Arg Ser Met Phe His Gly Cys Pro Thr Arg Asp Phe Ser Ala Ala Trp
 50 55 60
 50 gtc aaa gac ccg gaa acc gga gaa acc cgc ccc aag ctg cac aga gtt 240
 Val Lys Asp Pro Glu Thr Gly Glu Thr Arg Pro Lys Leu His Arg Val
 65 70 75 80
 ggc acc cgc tca ctt tcc cgg tgc cag tac gtt gcc ctg acc cac ccg 288

EP 1 076 094 A2

	Gly	Thr	Arg	Ser	Leu	Ser	Arg	Cys	Gln	Tyr	Val	Ala	Leu	Thr	His	Pro	
					85					90					95		
5	cag	cgc	tcc	gcg	gtg	ctg	gtc	tta	gac	atc	gac	atc	ccc	agc	cac	cag	336
	Gln	Arg	Ser	Ala	Val	Leu	Val	Leu	Asp	Ile	Asp	Ile	Pro	Ser	His	Gln	
					100				105						110		
	gcc	ggc	ggg	aac	atc	gag	cac	ctt	cac	ccg	cag	gta	tac	gcc	acc	ttg	384
	Ala	Gly	Gly	Asn	Ile	Glu	His	Leu	His	Pro	Gln	Val	Tyr	Ala	Thr	Leu	
					115				120						125		
10	gag	cgt	tgg	gca	cgg	gtg	gag	aaa	gcg	ccg	gcc	tgg	atc	ggg	gtg	aac	432
	Glu	Arg	Trp	Ala	Arg	Val	Glu	Lys	Ala	Pro	Ala	Trp	Ile	Gly	Val	Asn	
					130				135						140		
	ccg	ttg	tcg	gga	aag	tgc	cag	ctc	atc	tgg	tgc	att	gac	ccg	gtg	ttc	480
	Pro	Leu	Ser	Gly	Lys	Cys	Gln	Leu	Ile	Trp	Cys	Ile	Asp	Pro	Val	Phe	
					145				150						155		
15	gcc	gcc	gag	ggc	acc	agc	tcg	aac	acc	cgc	ctg	cta	gcg	gcc	acc		528
	Ala	Ala	Glu	Gly	Thr	Ser	Ser	Asn	Thr	Arg	Leu	Leu	Ala	Ala	Thr		
					165				170						175		
	acc	gag	gaa	atg	acc	cgt	gtg	ttc	ggc	gct	gac	cag	gca	ttt	tcc	cac	576
	Thr	Glu	Glu	Met	Thr	Arg	Val	Phe	Gly	Ala	Asp	Gln	Ala	Phe	Ser	His	
					180				185						190		
20	cgg	ctg	agc	cgg	tgg	ccg	ctg	cat	gtt	tct	gat	gat	ccg	acc	gcg	tac	624
	Arg	Leu	Ser	Arg	Trp	Pro	Leu	His	Val	Ser	Asp	Asp	Pro	Thr	Ala	Tyr	
					195				200						205		
	tcc	tgg	cac	tgc	cag	cac	aac	cga	gtc	gat	acg	ctt	gat	gag	ctg	atg	672
	Ser	Trp	His	Cys	Gln	His	Asn	Arg	Val	Asp	Thr	Leu	Asp	Glu	Leu	Met	
25					210				215						220		
	gag	gta	gcc	cgc	acg	atg	acc	gga	tca	aaa	aag	ccg	aga	aag	cac	gct	720
	Glu	Val	Ala	Arg	Thr	Met	Thr	Gly	Ser	Lys	Lys	Pro	Arg	Lys	His	Ala	
					225				230						235		
	cac	cag	gag	ttt	tcc	agc	ggt	cgg	gca	cgg	atc	gaa	gcc	gcg	cgg	aaa	768
	His	Gln	Glu	Phe	Ser	Ser	Gly	Arg	Ala	Arg	Ile	Glu	Ala	Ala	Arg	Lys	
30					245				250						255		
	gcc	acc	gca	gag	gcc	aaa	gcg	ctt	gcc	gcc	ttg	gac	gcc	acg	ctg	cct	816
	Ala	Thr	Ala	Glu	Ala	Lys	Ala	Leu	Ala	Ala	Leu	Asp	Ala	Thr	Leu	Pro	
					260				265						270		
35	acg	gcg	ctg	gag	gca	tca	ggc	gat	ctc	att	gac	ggg	gtg	cgg	gtg	ttg	864
	Thr	Ala	Leu	Glu	Ala	Ser	Gly	Asp	Leu	Ile	Asp	Gly	Val	Arg	Val	Leu	
					275				280						285		
	tgg	gca	gca	gag	ggg	cgt	gca	gcc	cgt	gat	gag	aca	gcg	ttt	cgc	cat	912
	Trp	Ala	Ala	Glu	Gly	Arg	Ala	Ala	Arg	Asp	Glu	Thr	Ala	Phe	Arg	His	
					290				295						300		
40	gcg	ttg	acc	gtg	ggt	tat	cag	ctt	aaa	gcc	gca	ggt	gaa	cgc	ctg	aaa	960
	Ala	Leu	Thr	Val	Gly	Tyr	Gln	Leu	Lys	Ala	Ala	Gly	Glu	Arg	Leu	Lys	
					305				310						315		
	gat	gcc	aag	atc	att	gat	gcg	tat	gag	cgt	gcc	tac	aac	gtc	gcc	cag	1008
	Asp	Ala	Lys	Ile	Ile	Asp	Ala	Tyr	Glu	Arg	Ala	Tyr	Asn	Val	Ala	Gln	
					325				330						335		
45	gcg	gtg	gga	gct	gat	ggg	cgt	gaa	ccg	gat	ctg	cct	gcc	atg	cgt	gat	1056
	Ala	Val	Gly	Ala	Asp	Gly	Arg	Glu	Pro	Asp	Leu	Pro	Ala	Met	Arg	Asp	
					340				345						350		
	cgt	cag	acg	atg	gcc	cgc	cgt	gtg	cgc	gcc	tac	gtc	gcc	aaa	ggc	cag	1104
	Arg	Gln	Thr	Met	Ala	Arg	Arg	Val	Arg	Ala	Tyr	Val	Ala	Lys	Gly	Gln	
					355				360						365		
50	ccc	acg	gtc	agc	gcc	agg	agc	aca	cag	acc	cag	agc	agt	cgg	ggc	cgg	1152
	Pro	Thr	Val	Ser	Ala	Arg	Ser	Thr	Gln	Thr	Gln	Ser	Ser	Arg	Gly	Arg	
					370				375						380		

55

aaa gcc ctg gcc acc atg ggc cgc aga ggc ggg caa aaa gcc gct gaa 1200
 Lys Ala Leu Ala Thr Met Gly Arg Arg Gly Gly Gln Lys Ala Ala Glu
 385 390 395 400
 5 cgc tgg aaa acc gat cct aac ggc aaa tac gcc caa gaa aac cgc caa 1248
 Arg Trp Lys Thr Asp Pro Asn Gly Lys Tyr Ala Gln Glu Asn Arg Gln
 405 410 415
 cga ctc gaa gct gca aac aag cga cgt caa gtc agc tgg aac aaa tac 1296
 Arg Leu Glu Ala Ala Asn Lys Arg Arg Gln Val Ser Trp Asn Lys Tyr
 420 425 430
 10 gcg agc acg aat tct ggc tac ggt ttc cga cac gta tgg gcc agc ttg 1344
 Ala Ser Thr Asn Ser Gly Tyr Gly Phe Arg His Val Trp Ala Ser Leu
 435 440 445
 gaa aaa tgc cta cgc gac gag caa atc atg gaa gaa aca ggg ctt tca 1392
 Glu Lys Cys Leu Arg Asp Glu Gln Ile Met Glu Glu Thr Gly Leu Ser
 450 455 460
 15 cga gct acc gtg acg cgc cat tgg gtg cac tgc gag agg ctg gcc tgc 1440
 Arg Ala Thr Val Thr Arg His Trp Val His Cys Glu Arg Leu Ala Cys
 465 470 475 480
 tgc caa atc ctt agg ggg gct cac gcc gta cac aga taa 1479
 Cys Gln Ile Leu Arg Gly Ala His Ala Val His Arg
 485 490

<210> 6

<211> 492

<212> PRT

25 <213> *Corynebacterium thermoaminogenes*

<400> 6

Met Thr Leu Ala Asp Ser Pro Gly Thr Tyr Thr Ala Asp Ala Trp Asn
 1 5 10 15
 30 Tyr Ser Thr Asp Leu Phe Asp Thr His Pro Glu Leu Ala Leu Arg Ser
 20 25 30
 Arg Gly Trp Asn His Gln Asp Ala Ala Glu Phe Leu Ala His Leu Asp
 35 40 45
 Arg Ser Met Phe His Gly Cys Pro Thr Arg Asp Phe Ser Ala Ala Trp
 50 55 60
 35 Val Lys Asp Pro Glu Thr Gly Glu Thr Arg Pro Lys Leu His Arg Val
 65 70 75 80
 Gly Thr Arg Ser Leu Ser Arg Cys Gln Tyr Val Ala Leu Thr His Pro
 85 90 95
 Gln Arg Ser Ala Val Leu Val Leu Asp Ile Asp Ile Pro Ser His Gln
 100 105 110
 40 Ala Gly Gly Asn Ile Glu His Leu His Pro Gln Val Tyr Ala Thr Leu
 115 120 125
 Glu Arg Trp Ala Arg Val Glu Lys Ala Pro Ala Trp Ile Gly Val Asn
 130 135 140
 Pro Leu Ser Gly Lys Cys Gln Leu Ile Trp Cys Ile Asp Pro Val Phe
 145 150 155 160
 45 Ala Ala Glu Gly Thr Thr Ser Ser Asn Thr Arg Leu Leu Ala Ala Thr
 165 170 175
 Thr Glu Glu Met Thr Arg Val Phe Gly Ala Asp Gln Ala Phe Ser His
 180 185 190
 50 Arg Leu Ser Arg Trp Pro Leu His Val Ser Asp Asp Pro Thr Ala Tyr
 195 200 205
 Ser Trp His Cys Gln His Asn Arg Val Asp Thr Leu Asp Glu Leu Met
 210 215 220

55

Glu Val Ala Arg Thr Met Thr Gly Ser Lys Lys Pro Arg Lys His Ala
 225 230 235 240
 His Gln Glu Phe Ser Ser Gly Arg Ala Arg Ile Glu Ala Ala Arg Lys
 245 250 255
 Ala Thr Ala Glu Ala Lys Ala Leu Ala Ala Leu Asp Ala Thr Leu Pro
 260 265 270
 Thr Ala Leu Glu Ala Ser Gly Asp Leu Ile Asp Gly Val Arg Val Leu
 275 280 285
 Trp Ala Ala Glu Gly Arg Ala Ala Arg Asp Glu Thr Ala Phe Arg His
 290 295 300
 Ala Leu Thr Val Gly Tyr Gln Leu Lys Ala Ala Gly Glu Arg Leu Lys
 305 310 315 320
 Asp Ala Lys Ile Ile Asp Ala Tyr Glu Arg Ala Tyr Asn Val Ala Gln
 325 330 335
 Ala Val Gly Ala Asp Gly Arg Glu Pro Asp Leu Pro Ala Met Arg Asp
 340 345 350
 Arg Gln Thr Met Ala Arg Arg Val Arg Ala Tyr Val Ala Lys Gly Gln
 355 360 365
 Pro Thr Val Ser Ala Arg Ser Thr Gln Thr Gln Ser Ser Arg Gly Arg
 370 375 380
 Lys Ala Leu Ala Thr Met Gly Arg Arg Gly Gly Gln Lys Ala Ala Glu
 385 390 395 400
 Arg Trp Lys Thr Asp Pro Asn Gly Lys Tyr Ala Gln Glu Asn Arg Gln
 405 410 415
 Arg Leu Glu Ala Ala Asn Lys Arg Arg Gln Val Ser Trp Asn Lys Tyr
 420 425 430
 Ala Ser Thr Asn Ser Gly Tyr Gly Phe Arg His Val Trp Ala Ser Leu
 435 440 445
 Glu Lys Cys Leu Arg Asp Glu Gln Ile Met Glu Glu Thr Gly Leu Ser
 450 455 460
 Arg Ala Thr Val Thr Arg His Trp Val His Cys Glu Arg Leu Ala Cys
 465 470 475 480
 Cys Gln Ile Leu Arg Gly Ala His Ala Val His Arg
 485 490

<210> 7
 <211> 1377
 <212> DNA
 <213> *Corynebacterium thermoaminogenes*

<220>
 <221> CDS
 <222> (1)..(1374)

<400> 7
 atg act cta gcg gat tcg cca gga aca tac aca gca gat gcg tgg aat 48
 Met Thr Leu Ala Asp Ser Pro Gly Thr Tyr Thr Ala Asp Ala Trp Asn
 1 5 10 15
 tac tcc aca gat ctg ttc gac acc cac cct gag ctg gct tta cgc tcc 96
 Tyr Ser Thr Asp Leu Phe Asp Thr His Pro Glu Leu Ala Leu Arg Ser
 20 25 30
 cgg ggt tgg aat cac cag gac gcc gcc gag ttc ctg gcc cac ctg gat 144
 Arg Gly Trp Asn His Gln Asp Ala Ala Glu Phe Leu Ala His Leu Asp
 35 40 45
 cgc agc atg ttt cac ggg tgc ccc acc cgg gat ttc tcc gcg gcc tgg 192
 Arg Ser Met Phe His Gly Cys Pro Thr Arg Asp Phe Ser Ala Ala Trp

50 55 60
 gtc aaa gac ccg gag acc gga gaa acc cgc cct aag ctg cac aga gtc 240
 Val Lys Asp Pro Glu Thr Gly Glu Thr Arg Pro Lys Leu His Arg Val
 5 65 70 75 80
 ggc acc cgg tgc ctt tcc cga tgc cag tac gtc gcg ctg acc cac ccg 288
 Gly Thr Arg Ser Leu Ser Arg Cys Gln Tyr Val Ala Leu Thr His Pro
 85 90 95
 cag cgc tcc gcg gtg ctg gtc tta gac atc gac atc ccc agc cac cag 336
 Gln Arg Ser Ala Val Leu Val Leu Asp Ile Asp Ile Pro Ser His Gln
 10 100 105 110
 gcc gcc ggg aac atc gag cac ctt cac ccg cag gtc tac gcc acc ttg 384
 Ala Gly Gly Asn Ile Glu His Leu His Pro Gln Val Tyr Ala Thr Leu
 115 120 125
 gag cgc tgg gca cgg gtg gag aaa gcg ccg gcc tgg atc ggg gtg aac 432
 Glu Arg Trp Ala Arg Val Glu Lys Ala Pro Ala Trp Ile Gly Val Asn
 15 130 135 140
 ccg ttg tca gga aag tgc cag ctc atc tgg tgc att gac ccg gtg ttc 480
 Pro Leu Ser Gly Lys Cys Gln Leu Ile Trp Cys Ile Asp Pro Val Phe
 145 150 155 160
 gcc gcc gag ggc acc acc agc ccg aac acc cgc ctg cta gcg gcc acc 528
 Ala Ala Glu Gly Thr Thr Ser Pro Asn Thr Arg Leu Leu Ala Ala Thr
 20 165 170 175
 acc gag gaa atg acc cgt atg ttc gcc gct gac cag gca ttt tcc cac 576
 Thr Glu Glu Met Thr Arg Met Phe Gly Ala Asp Gln Ala Phe Ser His
 180 185 190
 cgg ctg agc cgg tgg ccg ctg cat gta tct gat gat ccg acc gcg tac 624
 Arg Leu Ser Arg Trp Pro Leu His Val Ser Asp Asp Pro Thr Ala Tyr
 25 195 200 205
 tcc tgg cac tgc cag cac aac cga gtc gat acg ctt gct gag ctg atg 672
 Ser Trp His Cys Gln His Asn Arg Val Asp Thr Leu Ala Glu Leu Met
 210 215 220
 gag gta gcc ccg acg atg acc gga tca aaa aag cca gat agc act gct 720
 Glu Val Ala Arg Thr Met Thr Gly Ser Lys Lys Pro Asp Ser Thr Ala
 225 230 235 240
 cac cag gag ttt tcc agc ggt cgg gca cgg atc gaa gcc gcg agg aaa 768
 His Gln Glu Phe Ser Ser Gly Arg Ala Arg Ile Glu Ala Ala Arg Lys
 35 245 250 255
 gcc acc gca gaa gcc aaa gcg ctt gct gcc tta gac gcc acg ctg cct 816
 Ala Thr Ala Glu Ala Lys Ala Leu Ala Ala Leu Asp Ala Thr Leu Pro
 260 265 270
 acg gcg ctg gag gca tca gcc gat ctc att gac ggg gtg cgg gtg ctg 864
 Thr Ala Leu Glu Ala Ser Gly Asp Leu Ile Asp Gly Val Arg Val Leu
 40 275 280 285
 tgg gca gca gag ggg cgt gca gcc cgt gat gag acg gcg ttt cgc cat 912
 Trp Ala Ala Glu Gly Arg Ala Ala Arg Asp Glu Thr Ala Phe Arg His
 290 295 300
 gcg ttg acc gtg ggg tat cag ctt aaa gcc gca ggt gaa cgc ctg aaa 960
 Ala Leu Thr Val Gly Tyr Gln Leu Lys Ala Ala Gly Glu Arg Leu Lys
 45 305 310 315 320
 gac acc aag atc att gat gcg tat gag cgt gcc tac aac gtc gcc cag 1008
 Asp Thr Lys Ile Ile Asp Ala Tyr Glu Arg Ala Tyr Asn Val Ala Gln
 325 330 335
 gcg gtg ggg gct gat ggg cgt gag ccg gat ctg cct gcc atg cgt gat 1056
 Ala Val Gly Ala Asp Gly Arg Glu Pro Asp Leu Pro Ala Met Arg Asp
 50 340 345 350
 cgt cag acg ttg gcc cgt cgt gtg cgc gcc tac gtc gct aaa ggc cag ---1104

Arg Gln Thr Leu Ala Arg Arg Val Arg Ala Tyr Val Ala Lys Gly Gln
 355 360 365
 ccc acg gtg agc gcc agg agc aca cag acc cag agc agc cgg ggc agg 1152
 Pro Thr Val Ser Ala Arg Ser Thr Gln Thr Gln Ser Ser Arg Gly Arg
 370 375 380
 aaa gcc ctg gcc acc atg gga cgc aga ggc gca gcc acc tcg aat gca 1200
 Lys Ala Leu Ala Thr Met Gly Arg Arg Gly Ala Ala Thr Ser Asn Ala
 385 390 395 400
 cgc agg tgg gca gac cca gaa agc gat tac gcc cgc caa act cgg gag 1248
 Arg Arg Trp Ala Asp Pro Glu Ser Asp Tyr Ala Arg Gln Thr Arg Glu
 405 410 415
 cgt tta gcc cga gca atg agc ttc gta cat tca gca cag acg aga aca 1296
 Arg Leu Ala Arg Ala Met Ser Phe Val His Ser Ala Gln Thr Arg Thr
 420 425 430
 agg gcc gga tcc tgg cct acg ttt ccg agt gca agc gcc acg gtt acg 1344
 Arg Ala Gly Ser Trp Pro Thr Phe Pro Ser Ala Ser Ala Thr Val Thr
 435 440 445
 acc cca cga gca aag aag tcg caa cgg agc tag 1377
 Thr Pro Arg Ala Lys Lys Ser Gln Arg Ser
 450 455
 <210> 8
 <211> 458
 <212> PRT
 <213> Corynebacterium thermoaminogenes
 <400> 8
 Met Thr Leu Ala Asp Ser Pro Gly Thr Tyr Thr Ala Asp Ala Trp Asn
 1 5 10 15
 Tyr Ser Thr Asp Leu Phe Asp Thr His Pro Glu Leu Ala Leu Arg Ser
 20 25 30
 Arg Gly Trp Asn His Gln Asp Ala Ala Glu Phe Leu Ala His Leu Asp
 35 40 45
 Arg Ser Met Phe His Gly Cys Pro Thr Arg Asp Phe Ser Ala Ala Trp
 50 55 60
 Val Lys Asp Pro Glu Thr Gly Glu Thr Arg Pro Lys Leu His Arg Val
 65 70 75 80
 Gly Thr Arg Ser Leu Ser Arg Cys Gln Tyr Val Ala Leu Thr His Pro
 85 90 95
 Gln Arg Ser Ala Val Leu Val Leu Asp Ile Asp Ile Pro Ser His Gln
 100 105 110
 Ala Gly Gly Asn Ile Glu His Leu His Pro Gln Val Tyr Ala Thr Leu
 115 120 125
 Glu Arg Trp Ala Arg Val Glu Lys Ala Pro Ala Trp Ile Gly Val Asn
 130 135 140
 Pro Leu Ser Gly Lys Cys Gln Leu Ile Trp Cys Ile Asp Pro Val Phe
 145 150 155 160
 Ala Ala Glu Gly Thr Thr Ser Pro Asn Thr Arg Leu Leu Ala Ala Thr
 165 170 175
 Thr Glu Glu Met Thr Arg Met Phe Gly Ala Asp Gln Ala Phe Ser His
 180 185 190
 Arg Leu Ser Arg Trp Pro Leu His Val Ser Asp Asp Pro Thr Ala Tyr
 195 200 205
 Ser Trp His Cys Gln His Asn Arg Val Asp Thr Leu Ala Glu Leu Met
 210 215 220
 Glu Val Ala Arg Thr Met Thr Gly Ser Lys Lys Pro Asp Ser Thr Ala

225 His Gln Glu Phe Ser Ser Gly Arg Ala Arg Ile Glu Ala Ala Arg Lys 240
 245 250 255
 5 Ala Thr Ala Glu Ala Lys Ala Leu Ala Ala Leu Asp Ala Thr Leu Pro
 260 265 270
 Thr Ala Leu Glu Ala Ser Gly Asp Leu Ile Asp Gly Val Arg Val Leu
 275 280 285
 Trp Ala Ala Glu Gly Arg Ala Ala Arg Asp Glu Thr Ala Phe Arg His
 290 295 300
 10 Ala Leu Thr Val Gly Tyr Gln Leu Lys Ala Ala Gly Glu Arg Leu Lys
 305 310 315 320
 Asp Thr Lys Ile Ile Asp Ala Tyr Glu Arg Ala Tyr Asn Val Ala Gln
 325 330 335
 15 Ala Val Gly Ala Asp Gly Arg Glu Pro Asp Leu Pro Ala Met Arg Asp
 340 345 350
 Arg Gln Thr Leu Ala Arg Arg Val Arg Ala Tyr Val Ala Lys Gly Gln
 355 360 365
 Pro Thr Val Ser Ala Arg Ser Thr Gln Thr Gln Ser Ser Arg Gly Arg
 370 375 380
 20 Lys Ala Leu Ala Thr Met Gly Arg Arg Gly Ala Ala Thr Ser Asn Ala
 385 390 395 400
 Arg Arg Trp Ala Asp Pro Glu Ser Asp Tyr Ala Arg Gln Thr Arg Glu
 405 410 415
 Arg Leu Ala Arg Ala Met Ser Phe Val His Ser Ala Gln Thr Arg Thr
 420 425 430
 25 Arg Ala Gly Ser Trp Pro Thr Phe Pro Ser Ala Ser Ala Thr Val Thr
 435 440 445
 Thr Pro Arg Ala Lys Lys Ser Gln Arg Ser
 450 455

 <210> 9
 <211> 4369
 <212> DNA
 <213> *Corynebacterium thermoaminogenes*

 <220>
 <221> CDS
 35 <222> (1)..(1476)

 <400> 9
 atg act cta gcg gat tcg cca gga aca tac aca gca gat gcg tgg aat 48
 Met Thr Leu Ala Asp Ser Pro Gly Thr Tyr Thr Ala Asp Ala Trp Asn
 1 5 10 15
 40 tac tcc act gat ctg ttc gac acc cac cct gag ctg gct tta cgc tcc 96
 Tyr Ser Thr Asp Leu Phe Asp Thr His Pro Glu Leu Ala Leu Arg Ser
 20 25 30
 45 cgg ggt tgg aat cac cag gac gcc gca gag ttc ctg gcc cac ctg gat 144
 Arg Gly Trp Asn His Gln Asp Ala Ala Glu Phe Leu Ala His Leu Asp
 35 40 45
 cgc agc atg ttt cac ggg tgc ccc acc cgg gat ttc tcc gcg gcc tgg 192
 Arg Ser Met Phe His Gly Cys Pro Thr Arg Asp Phe Ser Ala Ala Trp
 50 55 60
 50 gtc aaa gac ccg gaa acc gga gaa acc cgc ccc aag ctg cac aga gtt 240
 Val Lys Asp Pro Glu Thr Gly Glu Thr Arg Pro Lys Leu His Arg Val
 65 70 75 80
 ggc acc cgc tca ctt tcc cgg tgc cag tac gtt gcc ctg acc cac ccg 288

	Gly	Thr	Arg	Ser	Leu	Ser	Arg	Cys	Gln	Tyr	Val	Ala	Leu	Thr	His	Pro	
				85						90					95		
5	cag	cgc	tcc	gcg	gtg	ctg	gtc	tta	gac	atc	gac	atc	ccc	agc	cac	cag	336
	Gln	Arg	Ser	Ala	Val	Leu	Val	Leu	Asp	Ile	Asp	Ile	Pro	Ser	His	Gln	
				100					105					110			
	gcc	ggc	ggg	aac	atc	gag	cac	ctt	cac	ccg	cag	gtg	tac	gcc	acc	ttg	384
	Ala	Gly	Gly	Asn	Ile	Glu	His	Leu	His	Pro	Gln	Val	Tyr	Ala	Thr	Leu	
				115				120					125				
10	gag	cgt	tgg	gca	cgg	gtg	gag	aaa	gcg	ccg	gcc	tgg	atc	ggg	gtg	aac	432
	Glu	Arg	Trp	Ala	Arg	Val	Glu	Lys	Ala	Pro	Ala	Trp	Ile	Gly	Val	Asn	
				130				135				140					
	ccg	tgt	tgc	gga	aag	tgc	cag	ctc	atc	tgg	tgc	att	gac	ccg	gtg	ttc	480
	Pro	Leu	Ser	Gly	Lys	Cys	Gln	Leu	Ile	Trp	Cys	Ile	Asp	Pro	Val	Phe	
				145			150				155				160		
15	gcc	ggc	gag	ggc	acc	acc	agc	tgc	aac	acc	cgc	ctg	cta	gcg	gcc	acc	528
	Ala	Ala	Glu	Gly	Thr	Thr	Ser	Ser	Asn	Thr	Arg	Leu	Leu	Ala	Ala	Thr	
				165						170			175				
	acc	gag	gaa	atg	acc	cgt	gtg	ttc	ggc	gct	gac	cag	gca	ttt	tcc	cac	576
	Thr	Glu	Glu	Met	Thr	Arg	Val	Phe	Gly	Ala	Asp	Gln	Ala	Phe	Ser	His	
				180					185				190				
20	cgg	ctg	agc	cgg	tgg	ccg	ctg	cat	gtt	ttt	gat	gat	ccg	acc	gcg	tac	624
	Arg	Leu	Ser	Arg	Trp	Pro	Leu	His	Val	Phe	Asp	Asp	Pro	Thr	Ala	Tyr	
				195				200					205				
	tcc	tgg	cac	tgc	cag	cac	aac	cga	gtc	gat	att	ctt	gat	gag	ctg	atg	672
	Ser	Trp	His	Cys	Gln	His	Asn	Arg	Val	Asp	Ile	Leu	Asp	Glu	Leu	Met	
				210			215					220					
25	gag	gta	gcc	cgc	acg	atg	acc	gga	tca	aaa	aag	ccg	aga	aag	cac	gct	720
	Glu	Val	Ala	Arg	Thr	Met	Thr	Gly	Ser	Lys	Lys	Pro	Arg	Lys	His	Ala	
				225			230				235				240		
	cac	cag	gag	ttt	tcc	agc	ggt	cgg	gca	cgg	atc	gaa	gcc	gcg	cgg	aaa	768
	His	Gln	Glu	Phe	Ser	Ser	Gly	Arg	Ala	Arg	Ile	Glu	Ala	Ala	Arg	Lys	
				245					250				255				
30	gcc	acc	gca	gag	gcc	aaa	gcg	ctt	gcc	gcc	ttg	gac	gcc	acg	ctg	cct	816
	Ala	Thr	Ala	Glu	Ala	Lys	Ala	Leu	Ala	Ala	Leu	Asp	Ala	Thr	Leu	Pro	
				260				265					270				
	acg	gcg	ctg	gag	gca	tca	ggc	gat	ctc	att	gac	ggg	gtg	cgg	gtg	ttg	864
	Thr	Ala	Leu	Glu	Ala	Ser	Gly	Asp	Leu	Ile	Asp	Gly	Val	Arg	Val	Leu	
				275			280						285				
	tgg	gca	gca	gag	ggg	cgt	gca	gcc	cgt	gat	gag	aca	gcg	ttt	cgc	cat	912
	Trp	Ala	Ala	Glu	Gly	Arg	Ala	Ala	Arg	Asp	Glu	Thr	Ala	Phe	Arg	His	
				290			295					300					
40	gcg	ttg	acc	gtg	ggt	tat	cag	ctt	aaa	gcc	gca	ggt	gaa	cgc	ctg	aaa	960
	Ala	Leu	Thr	Val	Gly	Tyr	Gln	Leu	Lys	Ala	Ala	Gly	Glu	Arg	Leu	Lys	
				305			310					315			320		
	gat	gcc	aag	atc	att	gat	gcg	tat	gag	cgt	gcc	tac	aac	gtc	gcc	cag	1008
	Asp	Ala	Lys	Ile	Ile	Asp	Ala	Tyr	Glu	Arg	Ala	Tyr	Asn	Val	Ala	Gln	
				325					330				335				
45	gcg	gtg	gga	gct	gat	ggg	cgt	gaa	ccg	gat	ctg	cct	gcc	atg	cgt	gat	1056
	Ala	Val	Gly	Ala	Asp	Gly	Arg	Glu	Pro	Asp	Leu	Pro	Ala	Met	Arg	Asp	
				340					345				350				
	cgt	cag	acg	atg	gcc	cgc	cgt	gtg	cgc	gcc	tac	gtc	gcc	aaa	ggc	cag	1104
	Arg	Gln	Thr	Met	Ala	Arg	Arg	Val	Arg	Ala	Tyr	Val	Ala	Lys	Gly	Gln	
				355				360					365				
50	ccc	acg	gtc	agc	gcc	agg	agc	aca	cag	acc	cag	agc	agt	cgg	ggc	cgg	1152
	Pro	Thr	Val	Ser	Ala	Arg	Ser	Thr	Gln	Thr	Gln	Ser	Ser	Arg	Gly	Arg	
				370				375					380				

aaa gcc ctg gcc acc atg ggc cgc aga ggc ggg caa aaa gcc gct gaa 1200
 Lys Ala Leu Ala Thr Met Gly Arg Arg Gly Gly Gln Lys Ala Ala Glu
 385 390 395 400
 5 cgc tgg aaa acc gat cct aac ggc aaa tac gcc caa gaa aac cgc caa 1248
 Arg Trp Lys Thr Asp Pro Asn Gly Lys Tyr Ala Gln Glu Asn Arg Gln
 405 410 415
 cga ctc gaa gct gca aac aag cga cgt caa gtc agc tgg aac aaa tac 1296
 Arg Leu Glu Ala Ala Asn Lys Arg Arg Gln Val Ser Trp Asn Lys Tyr
 420 425 430
 10 gcg agc acg aat tct ggc tac ggt ttc cga cac gta tgg gcc agc ttg 1344
 Ala Ser Thr Asn Ser Gly Tyr Gly Phe Arg His Val Trp Ala Ser Leu
 435 440 445
 gaa aaa tgc cta cgc gac gag caa atc atg gaa gaa aca ggg ctt tca 1392
 Glu Lys Cys Leu Arg Asp Glu Gln Ile Met Glu Glu Thr Gly Leu Ser
 450 455 460
 15 cga gct acc gtg acg cgc cat tgg gtg cac tgc gag agg ctg gcc tgc 1440
 Arg Ala Thr Val Thr Arg His Trp Val His Cys Glu Arg Leu Ala Cys
 465 470 475 480
 tgc caa atc ctt agg ggg gct cac gcc gta cac aga taacgggtcc cacc 1492
 Cys Gln Ile Leu Arg Gly Ala His Val His Arg
 485 490
 20 gtaggggtag cgttggtcc ctgaagctcc ggctcccatc cctctcagc actccctccc 1552
 cgaggggggg gctcacgcc tagacagata acggttccca ccccgtaggg gtagcgcttg 1612
 gtccctgaag ctctcacttc tggctccctt cctggccctc cttagtgcc caccataaa 1672
 tgcaaatgc cgtcagcaga caacggttcc caccctggg gtcctcaca caggctgcat 1732
 25 cagggtcttc gacgcttgct ggcttcatcc atcaactgct gggatgatc ctcgaacga 1792
 tccttgatcg cgaagtcttc gaaatcagcg gcagcttgct cccagtggat ccgttccagc 1852
 tccggcagcg ccgcccacag cgccatgcgc aagtaggttc cacgtgatcg cttagtgcgc 1912
 tcggcgagcg cgtccagcg ctcgatcagc tccggttcca ggcgcacgga gaccacgggg 1972
 ccgcgtcccg cggggttctt ctggttggtg gccatgagaa atttctctc gttcggtag 2032
 ttgtaaacaa tgtttacac gtgcgggga gagggtttt tattttctc ccgggcactt 2092
 30 tcgagacggg tcatgccgta agcagggcgc gtggccacac cgcactcggc gacgcaggtg 2152
 tcacttgctc cccgactgtc ggccgggaag gggcgcgca gagcgtccag gacgcggta 2212
 gagcgctgg gacggttctg gtggggactt ggtcgccca cggggcttta atcgcttaa 2272
 aacgcgcaca gcgcatctt tgccacgggc tagcgcgtga ccgctgcgc ctcacttgct 2332
 caggaagaaa atcattctc gcctaagcg cttcgcgcg tcgccctctc cgaaggggaa 2392
 35 aactaaccac acacctcatg cactaaagt ctgattgca ggtcagcgc ttttagcgtg 2452
 caaaaatagt gcgaaaacg gcgaaaatg gggcgcgaca atccctcag tggctccca 2512
 aaattcacct attcacatct gctactggct gacttcttc ccgacaagg gccctgtgag 2572
 ggcgcaggt gagccactt tacgtcccg agatccctt agggcgtatt cgagggtg 2632
 tcagtgaccc gcttcggcg ggtgggagta gccaaaagtc cgacatttt aacgaacgtt 2692
 cgttaaaatg ggggcagac tcagggacct ttgacctcag aaaccggcg aatcctgaat 2752
 40 gatcttgcg cagcagaccc tctcgatgtg gctgtccgg cagggagag tgcgcatgtt 2812
 ctctctcaag tcgtggagt tttagagcag atcgccgggt ctggggatag cgatttagac 2872
 gcggtgatg agcgtgatt gcagctcgat gcagacacgt tgaccttcat tgcccaggcg 2932
 ttggagggt tggcgacca ggcgagggc aaggatgcc tgaacgaat acgatatgt 2992
 gtgcccaatg cgggtgggaa atcccgcccc ggctgaccc ccgcgacgc agggcgaa 3052
 45 attgctcgga tgcttgctg gcggcagcga gcccggaacg cgcgcgccag cggcagccc 3112
 aggaggtcga agccgcgcgt ctccaggcg cactcgatct gaaaacccg caggagaccc 3172
 tggcagaggt agtccaggag cttcaggcca ccaccggat tatccgtgat cgaggggacg 3232
 tgccagcgtc gctgcgtccg ctggttaatg ctgcatccga actggtcaac gcagcgcaac 3292
 cggttgagga atctaagta ttcccaacc ggcgagtgcg tcgtgcagtt aaacgaaagt 3352
 50 ttgcgataag cgggtgatgt aactgatgga gatatttacc tgggggtgct tccagcgagg 3412
 ttgccaagtc cgatttggtt gaggattacc ccaaactgac ggggattatt caaaatccac 3472
 tgtccaaccg cttttccggt taccgcgct ccgatgcgc ctacgagaat agagcccatg 3532
 accattgcat tgtggctata tcccgcat 55

ccagcagcgc agcctgcaat gcgagcgcca atgataacca gggggagggc gcgaggcatt 3652
 actcgatttt catctgtggt ctgtcgctga atcgaagcag tgatggcttc ttcaaatgct 3712
 tcagggtttg acgtggggtc cgagactgtt gacgcagctt cctgcactgc cttgatgagg 3772
 5 acatcttcag ggatggaatc attgaacatt cctcccagct cagaagtgtt ttgaacgtta 3832
 gccgaaggga catgcacatc gggggaagcc tgggcggctg gagcaattaa agacagcgac 3892
 agtgaagcaa cgagagccgt tacagtggca cgagttttta aatacatgag gcgaacttaa 3952
 caaaccattg ataggttgtc gtgcggtaaa gataagaaaa ggataaagat atgaaaacgt 4012
 tatttatgaa tctcttaggt gccgcgcttg taggagcggg aatcatggtc ttgacatggt 4072
 10 tatttattga ttttgatgca cctggagcat ggctcggatt ctttattatc accaccatca 4132
 gtgattgctg ctttagaagt catccacgga ctttgggaaa aacggcaggg atcttccact 4192
 gacaatgatt gataaaacct ggttgaacgg aatacaaaac gcgcaaaata accaggcagt 4252
 taaaagaaaa accagataag ctgcaccaat acttgaaaaa tgttgaacgc cccgacagct 4312
 gtaactgtcg aggcgtcggc taacccccag tcatcagctg ggagaaagca ctcaaaa 4369

 15 <210> 10
 <211> 22
 <212> DNA
 <213> Artificial Sequence

 20 <220>
 <223> Description of Artificial Sequence: primer for
 amplifying replication origin of pYM2

 <400> 10
 aaccaggggg agggcgcgag gc 22

 25 <210> 11
 <211> 26
 <212> DNA
 <213> Artificial Sequence

 30 <220>
 <223> Description of Artificial Sequence: primer for
 amplifying replication origin of pYM2

 <400> 11
 35 tctcgtaggc tgcattccgag gcgggg 26

 <210> 12
 <211> 32
 <212> DNA
 <213> Artificial Sequence

 40 <220>
 <223> Description of Artificial Sequence: primer for
 amplifying replication origin of pYM2

 <400> 12
 45 gctctagagc aaccaggggg agggcgcgag gc 32

 <210> 13
 <211> 36
 <212> DNA
 50 <213> Artificial Sequence

 <220>

<223> Description of Artificial Sequence: primer for amplifying replication origin of pYM2

5 <400> 13
 gctctagagc tctcgtaggc tgcacggag gcgggg 36

10 <210> 14
 <211> 32
 <212> DNA
 <213> Artificial Sequence

<220>
<223> Description of Artificial Sequence: primer for
amplifying replication origin of pYM2

<400> 14
gctctagagc aaccaggggg agggcgcgag gc 32

20 <210> 15
 <211> 36
 <212> DNA
 <213> Artificial Sequence

25 <220>
<223> Description of Artificial Sequence: primer for
amplifying replication origin of pYM2

<400> 15
gctctagagc tctcgtaggc tgcattcggag gcgggg 36

30 <210> 16
 <211> 32
 <212> DNA
 <213> Artificial Sequence

35 <223> Description of Artificial Sequence:primer for
amplifying kanamycin resistant gene of
Streptococcus faecalis

40 <400> 16
cccgtaaact gcttgaacc caggacaata ac 32

<210> 17
<211> 30
<212> DNA
<213> Artificial Sequence

<223> Description of Artificial Sequence: primer for
amplifying kanamycin resistant gene of
Streptococcus faecalis

<400> 17
cccgtaaca tgtacttcag aaaagattag . - 30

<210> 18
 <211> 26
 <212> DNA
 <213> Artificial Sequence

<220>
 <223> Description of Artificial Sequence: primer for
 amplifying Escherichia coli cloning vector pHSG399

<400> 18
 gatattctacg tgccgatcaa cgtctc 26

<210> 19
 <211> 25
 <212> DNA
 <213> Artificial Sequence

<220>
 <223> Description of Artificial Sequence: primer for
 amplifying Escherichia coli cloning vector pHSG399

<400> 19
 aggccttttt ttaaggcagt tattg 25

30 Claims

1. A plasmid isolable from *Corynebacterium thermoaminogenes*, which comprises a gene coding for a Rep protein having the amino acid sequence shown in SEQ ID NO: 2 or an amino acid sequence having homology of 90% or more to the amino acid sequence shown in SEQ ID NO: 2, and has a size of about 4.4 kb or about 6 kb, or a derivative thereof.
2. The plasmid or the derivative thereof according to claim 1, which is isolable from *Corynebacterium thermoaminogenes* AJ12340 (FERM BP-1539), AJ12308 (FERM BP-1540) or AJ12310 (FERM BP-1542), has a size of about 4.4 kb and is represented by the restriction map shown in Fig. 1.
3. The plasmid or the derivative thereof according to claim 1, which is isolable from *Corynebacterium thermoaminogenes* AJ12309 (FERM BP-1541), has a size of about 6 kb and is represented by the restriction map shown in Fig. 2.
4. The plasmid or the derivative thereof according to claim 1, which comprises a gene coding for a Rep protein having the amino acid sequence shown in SEQ ID NO: 2, 4 or 6.
5. The plasmid or the derivative thereof according to claim 1, which comprises a gene coding for a Rep protein having the amino acid sequence shown in SEQ ID NO: 8.

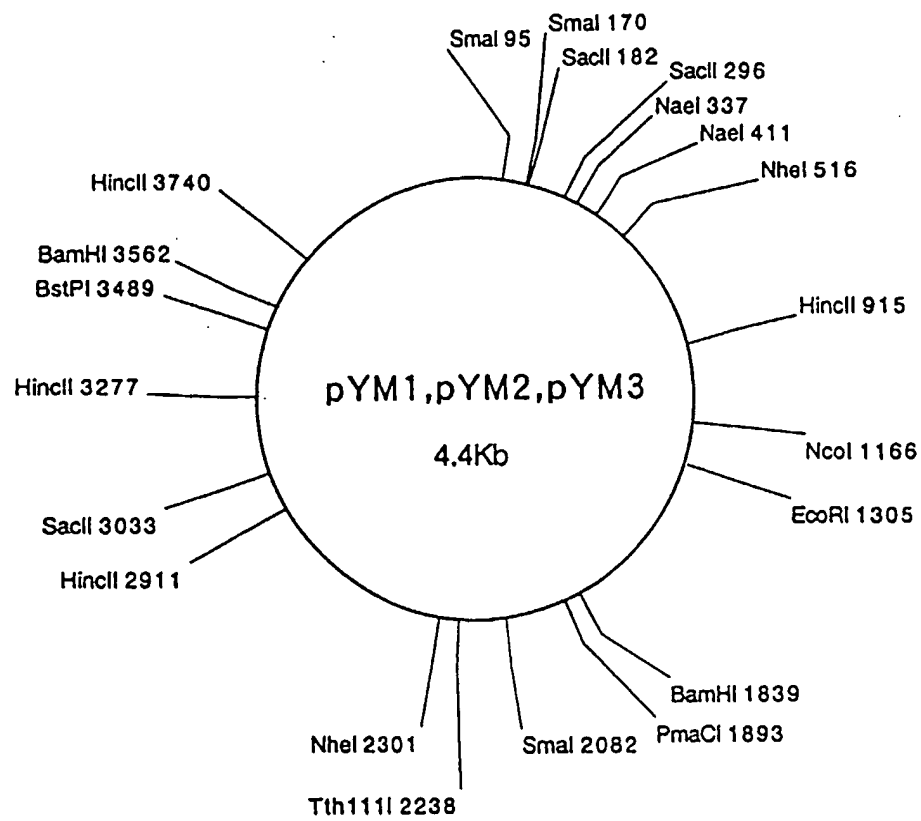


Fig. 1

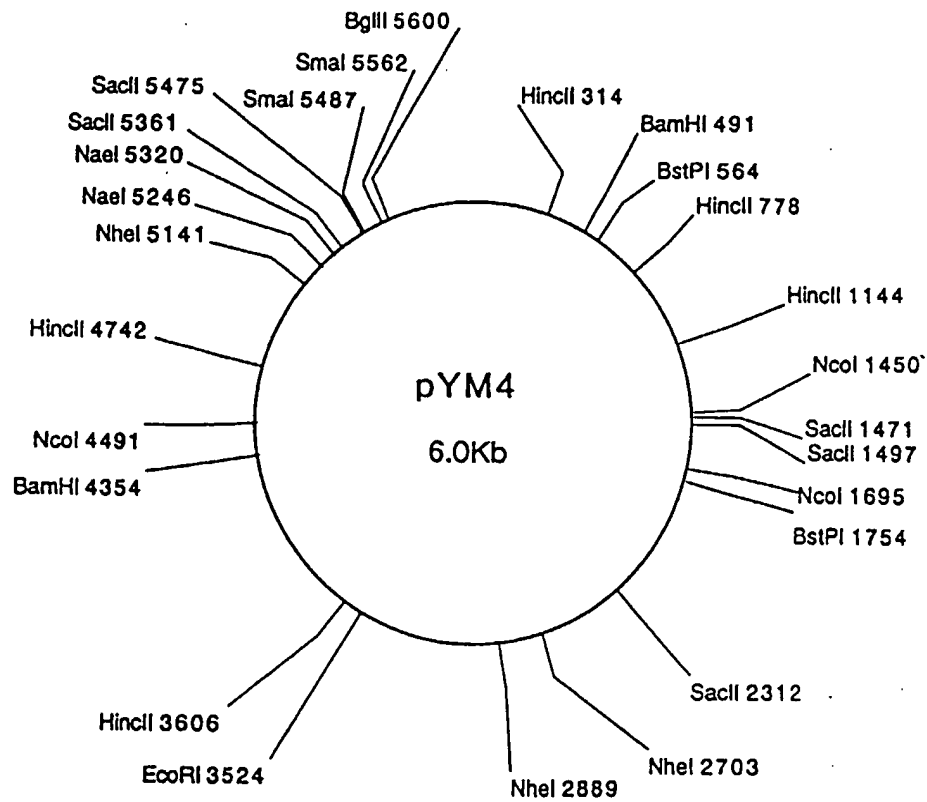


Fig. 2

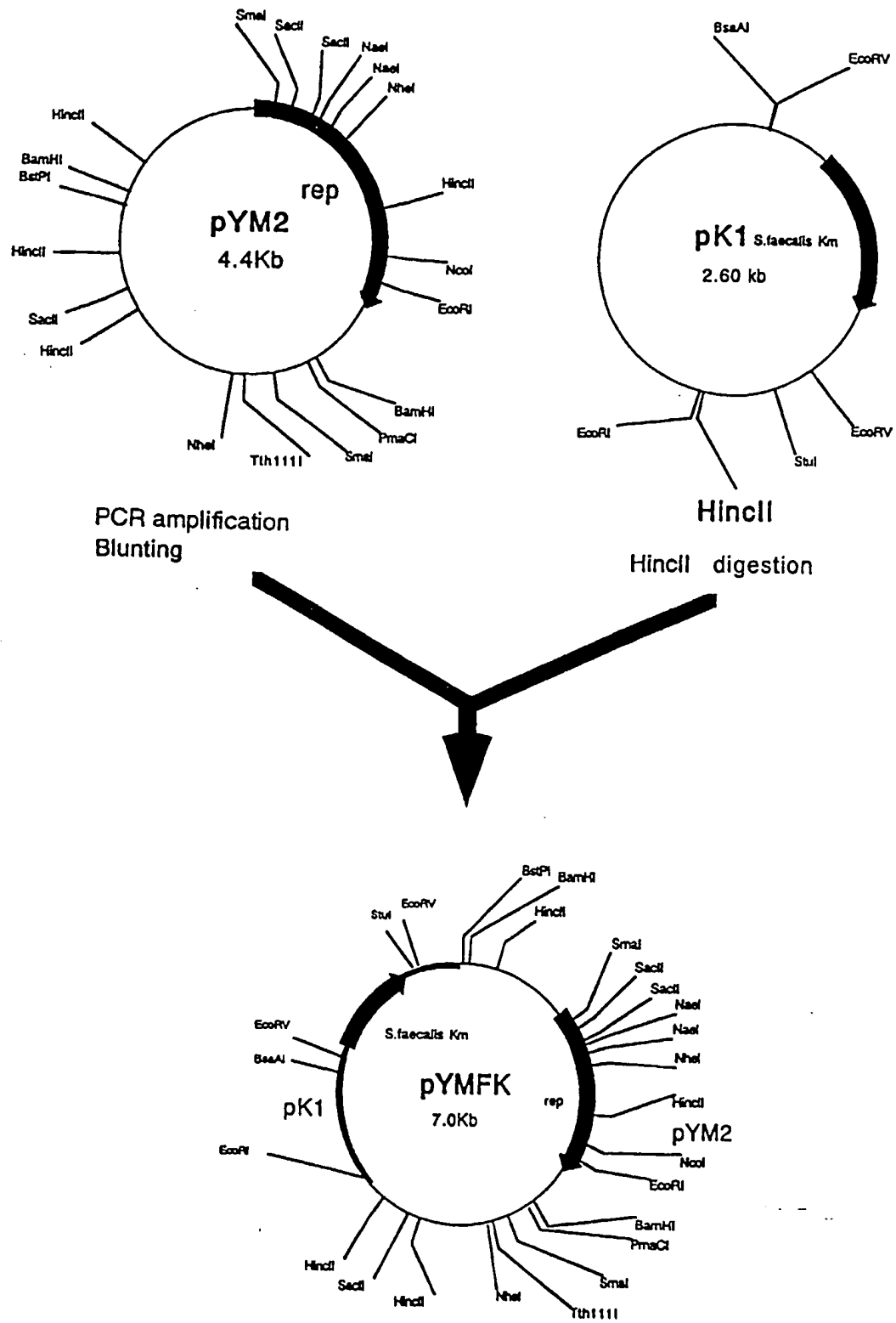


Fig. 3

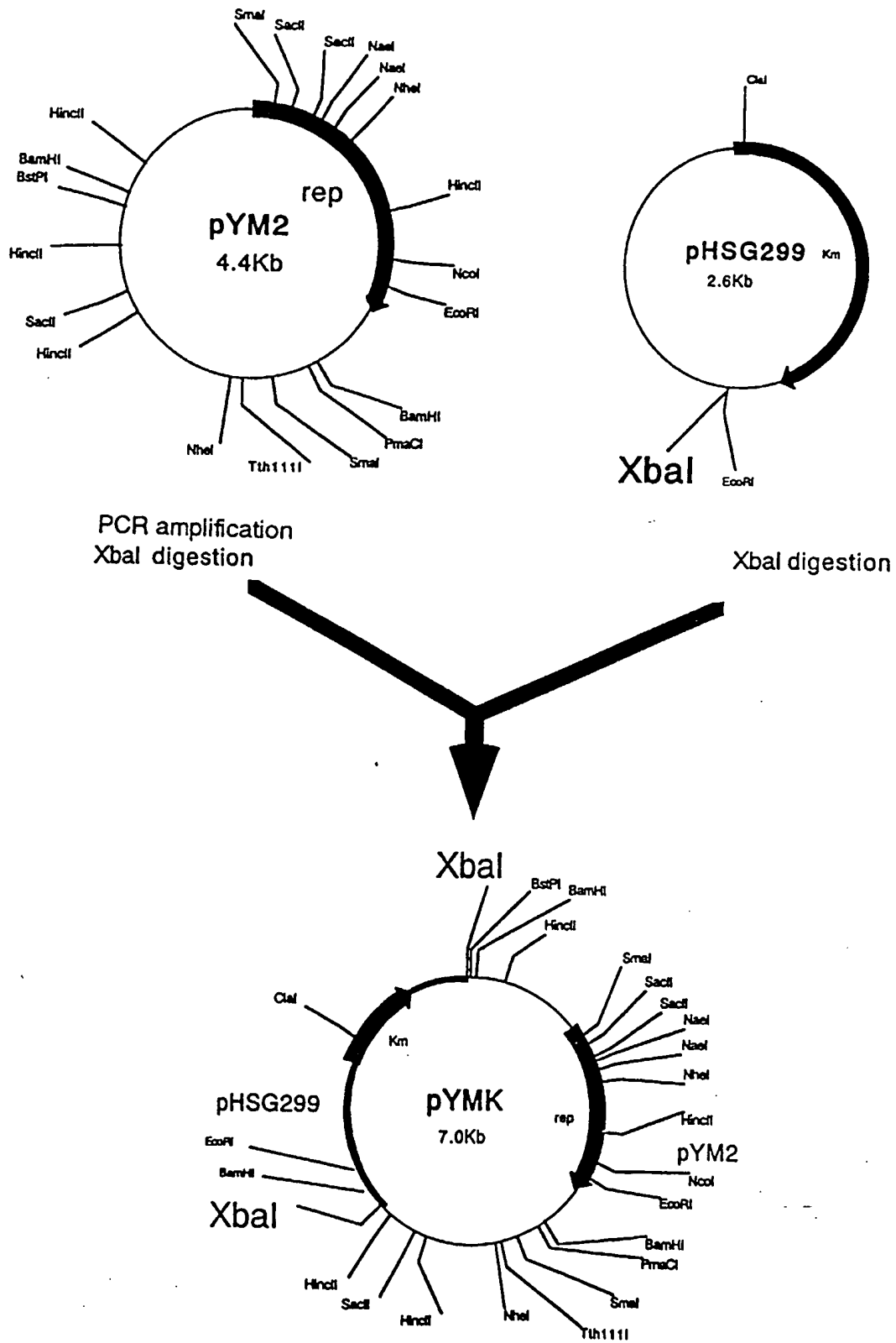


Fig. 4

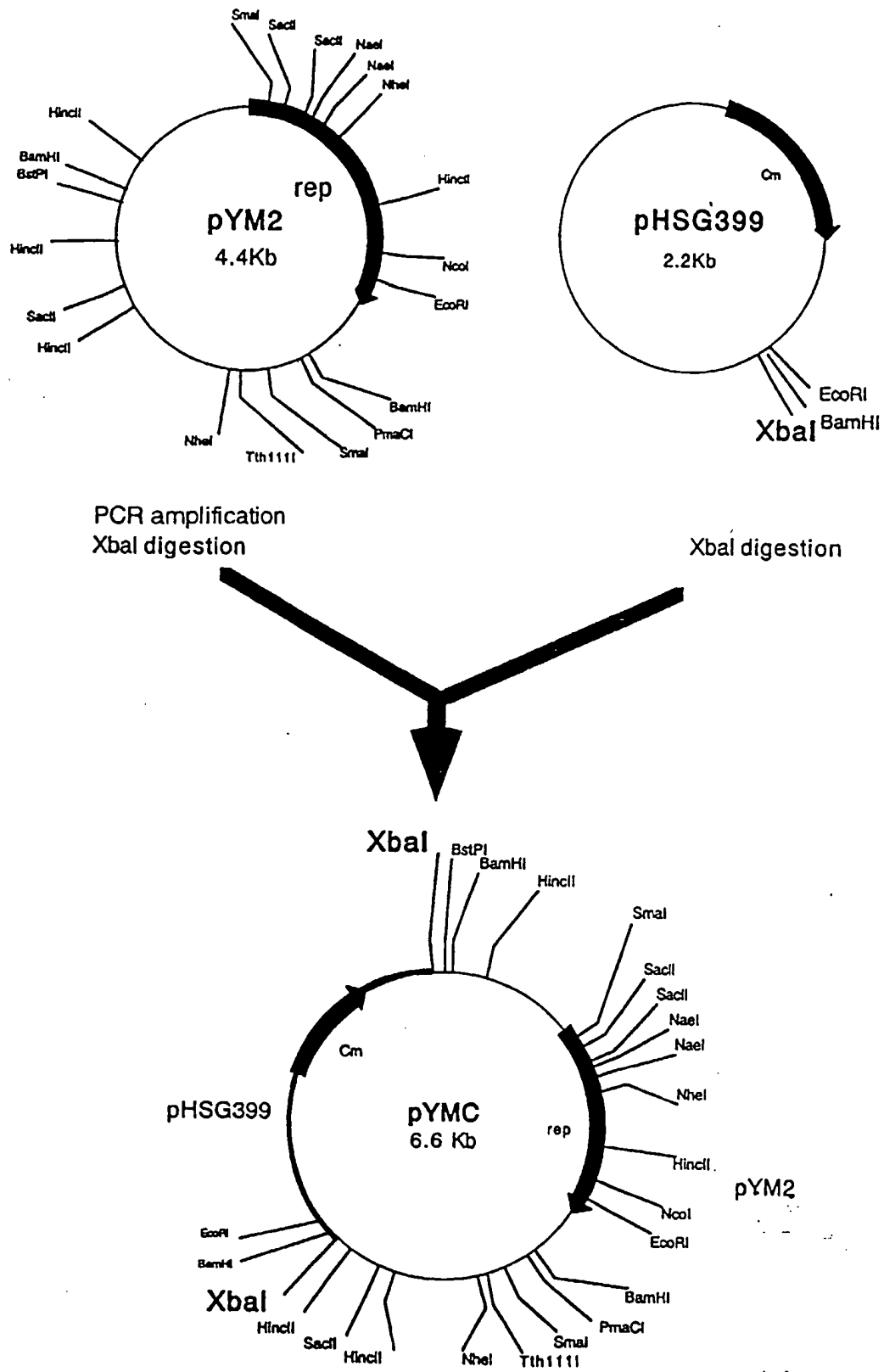


Fig. 5

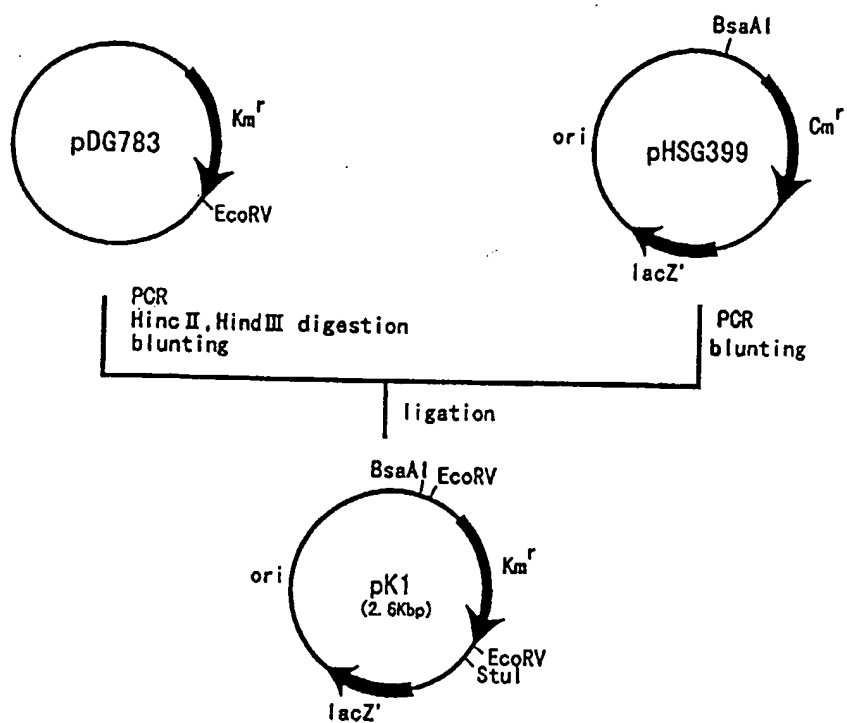
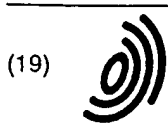


Fig. 6



(19)

Europäisches Patentamt

European Patent Office

Office européen des brevets



(11)

EP 1 076 094 A3

(12)

EUROPEAN PATENT APPLICATION

(88) Date of publication A3:

11.04.2001 Bulletin 2001/15

(51) Int. Cl.⁷: **C12N 15/74**

(43) Date of publication A2:

14.02.2001 Bulletin 2001/07

(21) Application number: **00117225.3**

(22) Date of filing: **11.08.2000**

(84) Designated Contracting States:

AT BE CH CY DE DK ES FI FR GB GR IE IT LI LU
MC NL PT SE

Designated Extension States:

AL LT LV MK RO SI

(30) Priority: **12.08.1999 JP 22839199**

(71) Applicant: **Ajinimoto Co., Inc.**

Tokyo 104 (JP)

(72) Inventors:

- Matsuzaki, Yumi,
c/o Ajinimoto Co., Inc.
Kawasaki-shi, Kanagawa (JP)
- Kimura, Eiichiro,
c/o Ajinimoto Co., Inc.
Kawasaki-shi, Kanagawa (JP)

- Nakamatsu, Tsuyoshi,
c/o Ajinimoto Co., Inc.
Kawasaki-shi, Kanagawa (JP)
- Kurahashi, Osamu,
c/o Ajinimoto Co., Inc.
Kawasaki-shi, Kanagawa (JP)
- Kawahara, Yoshio,
c/o Ajinimoto Co., Inc.
Kawasaki-shi, Kanagawa (JP)
- Sugimoto, Shinichi,
c/o Ajinimoto Co., Inc.
Kawasaki-shi, Kanagawa (JP)

(74) Representative: **HOFFMANN - EITLE**
Patent- und Rechtsanwälte
Arabellastrasse 4
81925 München (DE)

(54) **Plasmid capable of autonomous replication in coryneform bacteria**

(57) Plasmid isolated from *Corynebacterium thermoaminogenes* or a derivative thereof, wherein said plasmid has a size of about 4.4kb or about 6kb and comprises a gene coding for a Rep protein having the amino acid sequence shown in SEQ ID NO: 2 or a sequence at least 90% homologous to the same.

EP 1 076 094 A3



European Patent
Office

EUROPEAN SEARCH REPORT

Application Number
EP 00 11 7225

DOCUMENTS CONSIDERED TO BE RELEVANT			
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int.CI.7)
A	FR 2 612 937 A (AJINOMOTO KK) 30 September 1988 (1988-09-30) * claims 1,2 *	1-5	C12N15/74
A	EP 0 082 485 A (KYOWA HAKKO KOGYO KK) 29 June 1983 (1983-06-29) * page 6 - page 7, paragraph 2 * * examples 1.1,2 *	1-5	
A	EP 0 472 869 A (DEGUSSA) 4 March 1992 (1992-03-04) * page 5, paragraph 3 - last paragraph *	1-5	
			TECHNICAL FIELDS SEARCHED (Int.CI.7)
			C12N
The present search report has been drawn up for all claims			
Place of search THE HAGUE		Date of completion of the search 16 February 2001	Examiner Mata Vicente, T.
<p>CATEGORY OF CITED DOCUMENTS</p> <p>X : particularly relevant if taken alone Y : particularly relevant if combined with another document of the same category A : technological background O : non-written disclosure P : intermediate document</p> <p>T : theory or principle underlying the invention E : earlier patent document, but published on, or after the filing date D : document cited in the application L : document cited for other reasons & : member of the same patent family, corresponding document</p>			

EPO FORM 1603 03/92 (P04C01)

**ANNEX TO THE EUROPEAN SEARCH REPORT
ON EUROPEAN PATENT APPLICATION NO.**

EP 00 11 7225

This annex lists the patent family members relating to the patent documents cited in the above-mentioned European search report. The members are as contained in the European Patent Office EDP file on
The European Patent Office is in no way liable for these particulars which are merely given for the purpose of information.

16-02-2001

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
FR 2612937 A	30-09-1988	JP 2027856 C	26-02-1996
		JP 7063383 B	12-07-1995
		JP 63240779 A	06-10-1988
		AU 616168 B	24-10-1991
		AU 1161488 A	29-09-1988
		BR 8801289 A	25-10-1988
		KR 9606580 B	20-05-1996
		PH 25252 A	27-03-1991
		US 5250434 A	05-10-1993
EP 0082485 A	29-06-1983	JP 1960412 C	10-08-1995
		JP 6091827 B	16-11-1994
		JP 58105999 A	24-06-1983
		AT 37198 T	15-09-1988
		AU 556761 B	20-11-1986
		AU 9162982 A	23-06-1983
		CA 1199594 A	21-01-1986
		DE 3279030 D	20-10-1988
		DE 82485 T	22-12-1983
		ES 518348 D	16-04-1984
		ES 8404408 A	16-07-1984
		IL 67510 A	31-08-1988
		US 4710471 A	01-12-1987
EP 0472869 A	04-03-1992	DE 4027453 A	05-03-1992
		DE 59100582 D	16-12-1993
		DK 472869 T	03-01-1994
		JP 2603011 B	23-04-1997
		JP 4229183 A	18-08-1992
		US 5175108 A	29-12-1992

EPO FORM P0459

For more details about this annex : see Official Journal of the European Patent Office, No. 12/82